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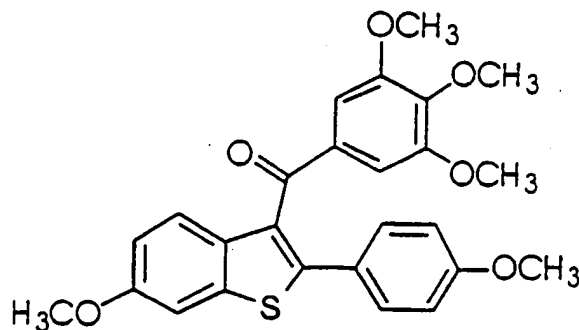
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(54) Title: INDOLE-CONTAINING AND COMBRETASTATIN-RELATED ANTI-MITOTIC AND ANTI-TUBULIN POLYMERIZATION AGENTS



(57) Abstract: Trimethoxyphenyl substituted indole ligands have been discovered which demonstrate impressive cytotoxicity as well as a remarkable ability to inhibit tubulin polymerization. Such compounds as well as related derivatives are excellent clinical candidates for the treatment of cancer in humans. In addition, certain of these ligands, as pro-drugs, may well prove to be tumor selective vascular targeting and destruction chemotherapeutic agents or to have anti-angiogenesis activity resulting in the selective prevention and/or destruction of tumor cell vasculature.

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INDOLE-CONTAINING AND COMBRETASTATIN-RELATED ANTI-MITOTIC AND ANTI-TUBULIN POLYMERIZATION AGENTS

BACKGROUND OF THE INVENTION

5 Tubulin is currently among the most attractive therapeutic targets in new drug design for the treatment of solid tumors.^{1c} The heralded success of vincristine and taxol along with the promise of combretastatin A-4 (CA-4) prodrug and dolastatin 10, to name just a few, have firmly established the clinical efficacy of these antimitotic agents for cancer treatment.

10 An aggressive chemotherapeutic strategy toward the treatment of solid-tumor cancers continues to rely on the development of architecturally new and biologically more potent anti-tumor, anti-mitotic agents which mediate their effect through a direct binding interaction with tubulin. A variety of clinically-promising compounds which demonstrate potent cytotoxicity and antitumor activity are known to effect their primary mode of action through an efficient inhibition of tubulin polymerization.¹ This class of compounds undergoes an initial interaction (binding) to the ubiquitous protein tubulin which in turn arrests the ability of tubulin to
15 polymerize into microtubules which are essential components for cell maintenance and division.² During metaphase of the cell cycle, the nuclear membrane is broken down and the cytoskeletal protein tubulin is able to form centrosomes (also called microtubule organizing centers) and through polymerization and depolymerization of tubulin the dividing chromosomes are separated. Currently, the most recognized and clinically useful members of this class of
20 antimitotic, antitumor agents are vinblastine and vincristine³ along with taxol.⁴ Additionally, the natural products rhizoxin,⁵ combretastatin A-4 and A-2,⁶ curacin A,¹ podophyllotoxin,⁷ epothilones A and B,⁸ dolastatin 10⁹ and welwistatin¹⁰ (to name just a few) as well as certain synthetic analogues including phenstatin,¹¹ the 2-styrylquinazolin-4(3H)-ones (SQO),¹² and highly oxygenated derivatives of *cis*- and *trans*-stilbene¹³ and dihydrostilbene are all known to
25 mediate their cytotoxic activity through a binding interaction with tubulin. The exact nature of this binding site interaction remains largely unknown, and definitely varies between the series of compounds. Photoaffinity labeling and other binding site elucidation techniques have identified several key binding sites on tubulin: colchicine site, vinca alkaloid site, and a site on the polymerized microtubule to which taxol binds.^{1a,14}

SUMMARY OF THE INVENTION

An important basic and essential aspect of this work requires a detailed understanding, on the molecular level, of the "small molecule" binding domain of both the α and β subunits of tubulin. The tertiary structure of the α , β tubulin heterodimer was reported earlier this year by
5 Downing and co-workers at a resolution of 3.7 Å using a technique known as electron crystallography.¹⁵ This brilliant accomplishment culminates decades of work directed toward the elucidation of this structure and should facilitate the identification of small molecule binding sites, such as the colchicine site, through techniques such as photoaffinity and chemical affinity labeling.

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BRIEF DESCRIPTION OF THE DRAWINGS

Figure 1 illustrates 3-(3', 4', 5' - trimethoxybenzoyl)-2-(4'-methoxyphenyl)-6-methoxybenzo[*b*]thiophene.

Figure 2 illustrates 2-(3', 4', 5'-trimethoxybenzoyl)-3-(4'-methoxyphenyl)-6-methoxybenzo[*b*]furan.

15

Figure 3 illustrates benzo[*b*]thiophene Phenol (BBT-OH).

Figure 4 illustrates benzo[*b*]thiophene prodrug (BBT-P).

Figure 5 illustrates *in vivo* biological data for benzo[*b*]thiophene prodrug (BBT-P).

Figure 6 illustrates a synthetic route for preparation of phenylindole derivatives.

Figure 7 illustrates a COSY NMR for 2-phenyl indole (aromatic region) a compound 31.

20

Figure 8 illustrates a cyclized isomer without aryl migration (no evidence for its formation).

Figure 9 illustrates a preparation of 2-phenylindole 31 in a one-pot reaction.

Figure 10 illustrates a designed synthetic route for preparation of indole-based analog.

Figure 11 illustrates a preparation of indole-based analog.

25

Figure 12 illustrates a synthesis of indole-based disodium prodrug salt.

Figure 13 illustrates another synthesis of indole-based disodium prodrug.

Figure 14 illustrates another synthesis of indole-based disodium prodrug.

Figure 15 illustrates a synthesis of indole based phosphoramidate prodrug.

Figure 16 illustrates another synthesis of indole-based disodium prodrug salt.

30

Figure 17A illustrates a combretastatin A-4 pro-drug.

Figure 17B illustrates a phosphoramidate analog 10.

Figure 18 illustrates a synthesis of phosphoramidate 10.

Figure 19 illustrates a model system used for phosphoramidate synthesis.

Figure 20 illustrates a synthesis of phosphoramidate 10 from from (Z)-3'-nitro combrestastatin analog 7B.

5 Figure 21 illustrates substituted 4-methoxyindole amines and/or phenols.

Figure 22 illustrates substituted 4-methoxyindole phosphate ester moieties and phosphoramidates.

Figure 23 illustrates further substituted 4-methoxyindole phosphate ester moieties and phosphoramidates.

10 Figure 24 illustrates substituted 6-methoxyindole amines and/or phenols.

Figure 25 illustrates substituted 6-methoxyindole phosphate ester moieties and phosphoramidates.

Figure 26 illustrates substituted 6-methoxyindole phosphate ester moieties and phosphoramidates.

15 Figure 27 illustrates substituted 4-methoxy-3-arylindole amines and/or phenols.

Figure 28 illustrates substituted 4-methoxy-3-arylindole phosphate moieties and phosphoramidates.

Figure 29 illustrates further substituted 4-methoxy-3-arylindole phosphate moieties and phosphoramidates.

20 Figure 30 illustrates 2-(4'-Methoxyphenyl)-3-(3'', 4'', 5''-trimethoxybenzoyl)-4-methoxyindole.

Figure 31 illustrates 2-(3', 4', 5'-Trimethoxybenzoyl)-3-(4''-methoxyphenyl)-6-methoxyindole.

25 Figure 32 illustrates 2-(3', 4', 5'-Trimethoxybenzoyl)-3-(4''-methoxyphenyl)-4-methoxyindole.

Figure 33 illustrates Disodium 2-(3'-phosphoramidate-4'-methoxyphenyl)-3-(3'', 4'', 5''-trimethoxybenzoyl)-6-methoxyindole.

Figure 34 illustrates 2-(3'-Hydroxy-4'-methoxyphenyl)-3-(3'', 4'', 5''-trimethoxybenzoyl)-4-methoxyindole.

30 Figure 35 illustrates 2-(3'-Amino-4'-methoxyphenyl)-3-(3'', 4'', 5''-trimethoxybenzoyl)-4-methoxyindole.

Figure 36 illustrates Disodium 2-[(4'-methoxyphenyl)-3'-O-phosphate]-3-(3'', 4'', 5''-trimethoxybenzoyl)-4-methoxyindole.

Figure 37 illustrates 2-(3'-Diethylphosphoramidate-4'-methoxyphenyl)-3-(3'', 4'', 5''-trimethoxybenzoyl)-4-methoxyindole.

5 Figure 38 illustrates Disodium 2-(3'-phosphoramidate-4'-methoxyphenyl)-3-(3'', 4'', 5''-trimethoxybenzoyl)-4-methoxyindole.

Figure 39 illustrates 2-(3',4',5'-trimethoxybenzoyl)-3-(3''-hydroxy-4''-methoxyphenyl)-6-methoxyindole.

10 Figure 40 illustrates 2-(3',4',5'-trimethoxybenzoyl)-3-(3''-amino-4''-methoxyphenyl)-6-methoxyindole.

Figure 41 illustrates Disodium 2-(3',4',5'-trimethoxybenzoyl)-3-[(4''-methoxyphenyl)-3''-O-phosphate]-6-methoxyindole.

Figure 42 illustrates 2-(3',4',5'-trimethoxybenzoyl)-3-[(4''-methoxyphenyl)-3''-diethylphosphoramidate]-6-methoxyindole.

15 Figure 43 illustrates Disodium 2-(3',4',5'-trimethoxybenzoyl)-3-[(4''-methoxyphenyl)-3''-phosphoramidate]-6-methoxyindole.

Figure 44 illustrates 2-(3',4',5'-trimethoxybenzoyl)-3-(3''-hydroxy-4''-methoxyphenyl)-4-methoxyindole.

20 Figure 45 illustrates 2-(3',4',5'-trimethoxybenzoyl)-3-(3''-amino-4''-methoxyphenyl)-4-methoxyindole.

Figure 46 illustrates Disodium 2-(3',4',5'-trimethoxybenzoyl)-3-[(4''-methoxyphenyl)-3''-O-phosphate]-4-methoxyindole.

Figure 47 illustrates 2-(3',4',5'-trimethoxybenzoyl)-3-[(4''-methoxyphenyl)-3''-diethylphosphoramidate]-4-methoxyindole.

25 Figure 48 illustrates Disodium 2-(3',4',5'-trimethoxybenzoyl)-3-[(4''-methoxyphenyl)-3''-phosphoramidate]-4-methoxyindole.

Figure 49 illustrates substituted 3-phosphoramidate derivatives of combretastatin A-4.

Figure 50 illustrates Disodium (Z)-1-[(4'-methoxyphenyl)-3'-phosphoramidate]-2-(3'',4'',5''-trimethoxyphenyl)ethene

30 Figure 51 illustrates substituted 3-phosphoramidate salts of combretastatin A-4.

DETAILED DESCRIPTION OF THE INVENTION

We have developed a working hypothesis suggesting that the discovery of new antimitotic agents may result from the judicious combination of a molecular template (scaffold) which in appropriately substituted form (ie. phenolic moieties, etc.) interacts with estrogen receptor (ER), suitably modified with structural features deemed imperative for tubulin binding (arylalkoxy groups, certain halogen substitutions, etc.). The methoxy aryl functionality seems especially important for increased interaction at the colchicine binding site in certain analogs.¹⁶ Upon formulation of this hypothesis concerning ER molecular templates, our initial design and synthesis efforts centered on benzo [*b*]thiophene ligands modeled after raloxifene, the selective estrogen receptor modulator (SERM) developed by Eli Lilly and Co.¹⁷ Our initial studies resulted in the preparation of a very active benzo[*b*]thiophene-based antitubulin agent.¹⁸⁻²¹ In further support of our hypothesis, recent studies have shown that certain estrogen receptor (ER) binding compounds as structurally modified estradiol congeners (2-methoxyestradiol, for example) interact with tubulin and inhibit tubulin polymerization.²² Estradiol is, of course, perhaps the most important estrogen in humans, and it is intriguing and instructive that the addition of the methoxy aryl motif to this compound makes it interactive with tubulin. It is also noteworthy that 2-methoxyestradiol is a natural mammalian metabolite of estradiol and may play a cell growth regulatory role especially prominent during pregnancy.

The design premise that molecular skeletons of traditional estrogen receptor (ER) binding compounds can be modified with structural motifs reminiscent of colchicine and combretastatin A-4 to produce inhibitors of tubulin polymerization has been validated by the benzo[*b*]thiophene and benzol[*b*]furan classes of new antimitotic agents.¹⁸⁻²¹ The lead compounds in each series (Figures 1 and 2), demonstrate remarkable biological activity against a variety of human cancer cell lines. For example, the 3,4,5-trimethoxybenzo[*b*]thiophene (Fig. 1) demonstrates potent cytotoxicity and inhibition of tubulin polymerization. In the NCI 60 cell line panel,²³ this compound produces a mean panel $GI_{50} = 2.63 \times 10^{-7}$ M (see Table I).

Inhibition of tubulin polymerization by 3-(3', 4', 5' - trimethoxybenzoyl)-2-(4'-methoxyphenyl)-6-methoxybenzo[*b*]thiophene. 50% inhibition of the maximum tubulin assembly rate with 1.1 μ M drug same assay with combretastatin A-4 gives a value of 0.73 μ M.

Human cancer cell line studies (*in vitro*) by 3-(3', 4', 5' - trimethoxybenzoyl)-2-(4'-methoxyphenyl)-6-methoxybenzo[*b*]thiophene.

Table I. Inhibition of tubulin polymerization by 2-(3', 4', 5'-trimethoxybenzoyl)-3-(4'-methoxyphenyl)-6-methoxybenzo[b]furan. IC₅₀ = 2.1 pM (totally flat at 4 pM).

Human cancer cell line studies (*in vitro*) by 2-(3', 4', 5'-trimethoxybenzoyl)-3-(4'-methoxyphenyl)-6-methoxybenzo[b]furan.

5	<u>Type of Cancer Cell Line</u>	<u>Cancer Cell Line</u>	<u>GI₅₀ (ug/mL)</u>
	Pancreas - adn	BXPC-3	0.03 8
	Neuroblast	SK-N-SH	0.025
	Thyroid ca	SW1736	0.047
	Lung-NSC	NCI-H460	0.041
10	Pharynx-sqam	FADU	0.03 5
	Prostate	DU-145	0.062

In addition, the phenolic derivative of the 3,4-5-trimethoxybenzo[b]thiophene compound (figure 3) has pronounced cytotoxicity and demonstrates outstanding inhibition of tubulin polymerization³⁶ and the pro-drug disodium phosphate salt form of this compound (Figure 4) demonstrates *in vitro* and *in vivo* cytotoxicity as a vascular targeting and destruction agent (which includes a component of tubulin binding (phenolic form of drug)^{36, 37} and subsequent inhibition of tubulin polymerization).

Initial *in vivo* studies are very encouraging (see Figure 5). Female scid mice were single dose ip administered with CA-4P, and benzo[b]thiophene phosphate prodrug at 400mg/kg (i.e. MDT of CA-4P) after one week of MHEC inoculation (1×10^6 /mouse). Studies were carried out through a collaboration with Professors Ronald W. Pero and Klaus Edvardsen, University of Lund, Sweden (Note: PbT Prodrug 20 is the same compound that is referred to as BBT-P).

Based on these promising research results, our interest in designing an indole based antimitotic agent was initiated, and a synthetic route (Schemes 1-4, see Figures 3A-D) was designed according to the synthesis of the benzo[b]thiophene derivatives.

The possibility clearly exists that some of the new indole-based ligands described herein, which are structurally related to combretastatin A-4, may also function through additional biological mechanisms involving anti-angiogenic activity. Clearly the ability to selectively disrupt the blood-flow to developing tumor cells is a potential breakthrough in the ever up-hill

battle against cancer. Certain phenylindoles have been noted for inhibiting tubulin polymerization.²⁷

A typical synthesis of indole-based ligand 33 is shown in Figures 6, 9, and 11. Secondary amine 30 was prepared by treatment of *m*-anisidine and 2-bromo-4-methoxyacetophenone under basic condition (ethanolic potassium hydroxide) at 0°C. Treatment of amine 30 with PPA resulted in the formation of two regioisomers. These isomers have poor solubility in EtOAc, CH₂Cl₂, and EtOH. Indole 31 was purified (from indole 32) by trituration in acetone. The structure of this isomer was confirmed by NMR analysis. COSY NMR was taken in order to study, in detail, the coupling relationship between the protons. The enlarged COSY spectrum for the aromatic region of ligand 31 is shown in Figure 5. This COSY NMR spectrum, shows a strong coupling between H^a and H^b which each appear as a doublet. H^c is coupled by the proton attached to the nitrogen into a small doublet. H^d is coupled only by H^c into a corresponding doublet, while H^e is coupled both by an ortho coupling (H^d) and by a meta coupling (H^f) into a doublet of doublet pattern. H^f is coupled by H^e into a doublet. Further evidence of the formation of 2-phenyl indole 31 is the chemical shift of the proton H^c on the ring which contains nitrogen. Though computer modeling (ChemDraw Ultra 4.5), the theoretical chemical shift value of 6.4 ppm is predicted for proton H^c (at the 3 position), which matches the peak shown in the actual NMR spectrum at 6.6 ppm. For the case where the proton is at the 2 position (Figure 8), the chemical shift is predicted to be 7.03 ppm, which does not match any peak in the spectrum that was obtained. Based collectively on these studies, the formation of isomer 31 is confirmed, and the migration of the methoxyphenyl system is evidenced. The other isomer (indole 32) is soluble in acetone and is much more difficult to obtain in pure form (see Figure 6).

Alternatively, another synthetic methodology can also be applied to the preparation of the desired 2-phenylindole. In 1984, Angerer and co-workers reported the synthesis of 2-phenylindoles in a one-pot reaction sequence (Figure 9) as a route toward the development of new therapeutic agents for the treatment of endocrine disorders.²⁵

Following this procedure (Figure 9), two arylindole regioisomers were obtained in good yield. Recrystallization in EtOH afforded the desired isomer, 2-phenylindole 31, as a white crystalline material.

In order to synthesize the indole-based analog 33, Friedel-Crafts acylation was carried out by treating indole 31 with 3,4,5-trimethoxybenzoyl chloride in the presence of the Lewis-Acid AlCl₃ (Figure 10). The reaction did not work under the regular conditions and only

starting material was obtained following work-up. Attempts to modify the reaction conditions by increasing the reaction temperature or using other Lewis Acids, such as TiCl_4 , proved futile as well. Starting material was recovered in all cases. One possible explanation for this result is the fact that the nitrogen atom (containing a lone pair of electrons and an acidic proton) may disrupt the acylation process. According to this analysis, a Grignard reagent (ethylmagnesium bromide) was used to protect this nitrogen prior to the Friedel-Crafts acylation step. Still, only starting material was obtained following the reaction. Therefore, a new synthetic approach was brought into this study.

In 1977, Inion and co-workers reported the synthesis of a variety of aminoalkoxy-4-benzoyl-3-indoles.²⁶ The benzoate indole product was prepared by treatment of indole with the appropriate benzoyl chloride with heating (130-150°C). HCl is generated under these conditions. A similar synthetic approach was used in the synthesis of the desired trimethoxybenzoate indole ligand 33 (Figure 11).

The precursor, indole 31, was mixed with trimethoxybenzoyl chloride. Since both reagents are solid, a solvent with a high boiling point was needed. 1,2-dichlorobenzene was chosen in this case since it has a boiling point of 180°C. Under these conditions, indole 33 was obtained in moderate yield following purification by flash column chromatography and recrystallization. NMR spectroscopy suggests that the structure of indole 33 is that indicated in Figure 11.

Based on promising results obtained with benzo[*b*]thiophene and benzofuran analogs, the preparation of phosphate salts is detailed in Figures 12-14, the preparation of analogs is detailed in Figures 15-16 and the preparation of similar indole-based phosphate prodrug salts and phosphoramidate derivatives is detailed in Figures 21-51.

In addition to the phosphate ester prodrugs that are described in this application for indole-based anti-mitotic agents, we have also discovered that phosphorous based prodrug derivatives of the nitrogen analog of combretastatin A-4 (CA-4) may have therapeutic advantages as selective tumor vasculature destruction agents. These compounds are primarily phosphoramidate derivatives and related phosphate dianions that are assembled on the 3-amino substituent of the nitrogen analog of CA-4. Although we describe two specific compounds and several obvious analogs, it should be apparent to anyone skilled in the art, that there are numerous other nitrogen phosphorous bond designs that might be assembled from the

3-amino-combretastatin A-4 structure and that would display similar functionality as prodrugs for the selective destruction of tumor vasculature.

Further significance is given to new drugs that bind to the colchicine site since it has recently been shown that combretastatin CA-4 also demonstrates anti-angiogenesis activity.²⁴ An emerging area of cancer chemotherapy centers on the development of both anti-angiogenesis drugs which disrupt the new microvessel formation of developing tumors and vascular targeting and destruction agents which selectively target the vasculature of tumor cells while leaving healthy cells intact. Combretastatin CA-4P prodrug (Figure 17A) is one of the leading new candidates from among a relatively small collection of known world compounds which display this vascular targeting. Discovered by Professor George R. Pettit (Arizona State University) from a willow tree (*combretum caffrum*) in South Africa in the 1970s, this compound is currently undergoing phase I clinical evaluation sponsored and licensed by OXiGENE, Inc.

Combretastatin A-4 (CA-4) is a potent inhibitor of tubulin polymerization which binds to the colchicine site on β -tubulin. Interestingly, CA-4 itself does not demonstrate destruction of tumor vasculature, while CA-4 prodrug is very active in terms of tumor vasculature destruction. It is very likely that the phosphate ester portion of the prodrug undergoes dephosphorylation (perhaps through the action of endothelial alkaline phosphatases) selectively at sites of enhanced vascularization to reveal the potent CA-4 itself which destroys the tumor cell through an inhibition of tubulin polymerization. The dephosphorylation event takes place selectively at tumor cells since tumor cells represent sites of prolific vascularization and alkaline phosphatases appear to be present at elevated concentrations in the endothelial cells lining tumor vasculature. This need for enhanced vascularization is not necessary for healthy cells. Hence, this dual-mode reactivity profile is clearly important in order to target tumor cells selectively over healthy cells. This is a proposal which has been advanced by Professor Ronald Pero (OXiGENE, Inc., University of Lund) for which a variety of strong evidence has been obtained.

Based in part on the good and promising biological results obtained for the 3'-nitrogen analogs of combretastatin A-4, a phosphoramidate analog has been prepared as a new combretastatin A-4 nitrogen prodrug (Figure 17B).

Phosphoramidate 10 below was obtained following the procedure reported by Taylor and coworkers for unrelated aryl amines.²⁸ Treatment of arylamine 7B with diethylchlorophosphite in anhydrous ether followed by oxidation with m-CPBA produced the phosphoramidate 10 in moderate yield (Figure 18).

A previous attempt in the synthesis of the phosphoramidate analog 10 utilized the methodology reported by Bilha Fisher and Larisa Sheihet.²⁹ This methodology presents a phosphoramidate intermediate, which can be isolated from the reduction of nitro aryl compounds to the corresponding aryl amines using diethylchlorophosphite as a biphilic reagent. The (Z)-nitro combretastatin analog 7B was considered a viable starting material for the synthesis of the phosphoramidate prodrug 10. This reaction was also tried using (Z)-1-(3',4',5'-trimethoxyphenyl)-2-(4"-nitrophenyl)ethene (synthesized in a similar manner as the other combretastatin containing analogs reported previously) as a model system (Figure 19). In neither case was the phosphoramidate product observed. It is thought that the presence of methoxy groups as strong electron donating substituents on the stilbene system disfavors the reaction (Figure 20).

It should be obvious to anyone skilled in the art of phosphate or phosphoramidate chemistry that there are numerous other synthetic methods which can be employed to prepare phosphoramidates (such as 10) and their related salts ($-\text{NHPO}_3^{-2}\text{Na}^+$).

Table II.³⁰ *In vitro* Human Cancer Cell Line Study of Phosphoramidate Analog 10. GI_{50} , TGI, and LC_{50} are reported as concentrations in $\mu\text{g/mL}$ ND = Not determined

Cell Type	Cell Line	GI_{50}	TGI	LC_{50}
Pancreas-a	BXPC-3	1.5×10^{-1}	5.7×10^{-1}	>10
Ovarian	OVCAR-3	1.9×10^{-1}	8.6×10^{-1}	>10
CNS	SF-295	2.4×10^{-1}	>10	>10
Lung-NSC	NCI-H460	3.5×10^{-1}	>10	>10
Colon	KM20L2	2.8×10^{-1}	6.1×10^{-1}	>10
Prostate	-DU-145	2.6×10^{-1}	2.6×10^{-1}	>10
Leukemia	P388	3.1×10^{-1}	ND	ND

Biological evaluation (*in vitro*) suggests that the phosphoramidate prodrug 10 is less effective than the corresponding amine 8 (Table II). Pettit and co-workers reported a similar loss in biological activity *in vitro* for the phosphate prodrugs of combretastatin A-4 and phenstatin compared to the original compounds (Table III).³¹ These results might be explained by the bulkiness of the phosphorous group and its steric hindrance toward binding site recognition. In

fact, Pettit and co-workers reported no inhibition of tubulin polymerization with the combretastatin prodrug while only a 40% activity is present for the phenstatin prodrug compared to phenstatin. The IC_{50} values for inhibition of tubulin polymerization are $1.2 \pm 0.1 \mu M$ for CA-4, $>80 \mu M$ for CA-4 prodrug, $1.0 \pm 0.2 \mu M$ for phenstatin and $21 \pm 3 \mu M$ for phenstatin prodrug; similar results are expected for the amino-CA-4 8 and the phosphoramidate 10.³¹ The IC_{50} for the amino-CA-4 8 is $1.2 \pm 0.02 \mu M$, and the phosphoramidate 10 has little if any activity.³²

Table III. Comparative GI_{50} Values Against Human Cancer Cell Lines for Amine-CA-4 8, Amine-CA-4 Prodrug 10, Phenstatin, Phenstatin Prodrug and Combretastatin A-4 Prodrug. GI_{50} values are reported as concentrations in $\mu g/mL$ ND = Not determined.^a Data obtained in collaboration with Dr. George R. Pettit.^{30b} Data obtained from synthesis of phenstatin phosphate.

Cell Type	Cell-Line	Amine-CA-4 8 ^a	Amine-CA-4 Prodrug 10 ^a	Phenstatin	Phenstatin Prodrug ^b	Combretastatin A-4 Prodrug ^b
Ovarian	OVCAR-3	ND	1.9×10^{-1}	2.3×10^{-3}	2.5×10^{-3}	2.3×10^{-2}
CNS	SF-295	ND	2.4×10^{-1}	5.2×10^{-2}	1.2×10^{-2}	3.6×10^{-2}
Lung-NSC	NCI-H460	6.8×10^{-4}	3.5×10^{-1}	5.7×10^{-3}	3.5×10^{-2}	2.9×10^{-2}
Colon	KM20L2	ND	2.8×10^{-1}	4.0×10^{-4}	2.7×10^{-1}	3.4×10^{-1}

In terms of *in vivo* systems, phosphoramidate analog 10 is able to provide a more soluble compound than the amine 8, thereby incrementing its bioavailability. Under, *in vivo* biological conditions, the P-N bond can be broken by serum phosphatases releasing the amine which can inhibit tubulin polymerization in a manner analogous to combretastatin

20 Anti-Angiogenesis

The growth of a tumor depends on the generation of blood vessels which will provide all the metabolites required during cell division. The development of anti-angiogenic compounds is especially useful in the treatment of solid tumors, since these compounds have the potential capability of selectively disrupting the vasculature of tumor cells while leaving healthy cells in a viable situation. The combretastatin A-4 prodrug has demonstrated anti-angiogenic activity since small doses of the drug are toxic to tumor vasculature.³⁴ Enhanced cytotoxic activity was observed against endothelial cells associated with the tumor vasculature of cancerous cells, while at the same time it was reported to have no effect against other endothelial cells which are

located distant from the tumor itself.^{34, 35} The mechanism of action of combretastatin A-4 prodrug, as an anti-angiogenic drug for cancer treatment, is under investigation because the development of blood vessels is crucial for the survival and growth of solid tumors. One proposed mechanism for anti-angiogenesis involves induction of apoptosis (cell suicide) of the cells instead of necrosis. An evaluation of the ability of the new phosphoramidate 10, along with structurally similar compounds, to induce apoptosis of endothelial cells will be undertaken in the near future.

Synthesis of the Phosphoramidate Analog

(Z)-1-(3'-Diethylphosphoramidate-4'-methoxyphenyl)-2-(3'', 4'', 5''-trimethoxyphenyl)ethene 10.

Diethylchlorophosphite (0.103 g, 0.66 mmol) was dissolved in anhydrous diethyl ether (2.5 ml) and cooled to -78°C. Diisopropylethyl amine (0.187 g, 1.45 mmol) was dissolved in Et₂O (1.0 ml) and added slowly over a period of 2 min to the reaction mixture by syringe. Amino-stilbene 8 was dissolved in Et₂O (1.0 mL) and added slowly to the reaction mixture by syringe. The reaction mixture was stirred under nitrogen at -78°C for 2 hours, followed by stirring for 1 hour at room temperature. The mixture was filtered, and the solvent was removed under reduced pressure. A yellow oil was obtained which was dissolved in dry CH₂Cl₂ (5 mL). The oil was cooled to -40°C and a solution of *m*-CPBA (0.193 g, 1.12 mmol) in CH₂Cl₂ (5 mL) was added. It was stirred over one hour at room temperature. After this time, the reaction mixture was cooled to -40°C and filtered through a sintered glass funnel. The liquid was collected with vigorous stirring over sodium sulfite (5%) (20 ml) in order to quench the reaction. The product was isolated by extraction with CH₂Cl₂, and washed with a saturated solution of NaHCO₃. The yellow oil which was obtained was dried over MgSO₄. Purification by flash chromatography (70/30, hexanes/EtOAc) afforded the phosphoramidate 10 as a yellow oil (0.130 g, 0.29 mmol, 44%).

¹H-NMR (CDCl₃, 360 MHz) δ 7.12 (d, *J* = 1.9 Hz, 1H, ArH), 6.88 (dd, *J* = 8.4 Hz, 2.0 Hz, 1H, ArH), 6.72 (dd, *J* = 8.4 Hz, 1.7, 1H, ArH), 6.49 (s, 2H, ArH), 6.51 (d, *J* = 12.1 Hz, 1H, vinyl CH), 6.41 (d, *J* = 12.1 Hz, 1H, vinyl CH), 5.67 (d, *J* = 10.0 Hz, NH), 4.02 (m, 4H, CH₂), 3.83 (s, 3 H, OCH₃), 3.83 (s, 3 H, OC₃), 3.68 (s, 6 H, OCH₃), 1.25 (t, 6 H, *J* = 7.1 Hz, CH₃).

^{13}C -NMR (CDCl_3 , 90 MHz) δ 152.7, 146.7, 146.6, 137.0, 132.7, 130.3, 129.7, 129.1, 129.0, 122.0, 117.0, 109.9, 106.0, 62.8, 60.7, 55.7, 16.1.

^{31}P -NMR (CDCl_3 , 145 MHz) δ 0.84.

HRMS (EI) M^+ calcd for $\text{C}_{22}\text{H}_{30}\text{NO}_7\text{P}$ 451.1760, found 451.1765.

5 Example 1

SYNTHESIS OF THE INDOLE-BASED ANTI-TUBULIN AGENTS

Preparation of 2-Phenyl Indole 31

Method I (2 steps):

10 To a well-stirred solution of KOH (0.926 g, 16.5 mmol) in EtOH (18 ml) and H_2O (9 ml) at rt was added *m*-anisidine (2.192 g, 17.80 mmol) by syringe. The solution was then stirred at 0°C . After 10 min, the solution of 2-bromo-4-methoxyacetophenone (4.09 g, 17.80 mmol) was added dropwise with an addition funnel over a 40 minute period. After 24 h, 0°C to rt, water was added. The product was isolated by extraction (1 H HCl, NaHCO_3 , brine, MgSO_4). The product was purified by recrystallization (50:50 EtOAc:hexanes) to afford secondary amine 30 (2.46 g, 9.07 mmol, 52%) as yellow solid.

^1H NMR (CDCl_3): δ 7.98 (2H, d, $J = 8.9$ Hz), 7.12 (1H, t, $J = 8.1$ Hz), 6.97 (2H, d, $J = 8.9$ Hz), 6.30 (3H, m), 4.54 (2H, s), 3.88 (3H, s), 3.79 (3H, s).

20 Polyphosphoric acid (PPA) was charged to a round-bottom flask and the temperature was raised to 80°C with vigorous stirring. To this flask was added the foregoing amine 30 (4.0 g, 14.74 mmol) in 6 portions over a 30 minute period. After 2 h, 80°C to 90°C , water was added. The product was isolated by extraction (EtOAc, NaHCO_3 , brine, MgSO_4). Purification by recrystallization (acetone) afforded indole 31 (0.544 g, 2.15 mmol, 15%) as a pale yellow solid.

25 ^1H NMR (CDCl_3): δ 11.24 (1H, br, s), 7.72 (2H, d, $J = 8.82$ Hz), 7.36 (1H, d, $J = 8.57$ Hz), 7.00 (2H, d, $J = 8.84$ Hz), 6.85 (1H, d, $J = 2.07$ Hz), 6.66 (1H, d, $J = 1.66$ Hz), 6.63 (1H, dd, $J = 8.59, 2.28$ Hz), 3.78 (3H, s), 3.77 (3H, s).

^{13}C NMR (CDCl_3): δ 158.15, 155.22, 137.44, 136.33, 125.60, 124.93, 122.82, 120.04, 114.07, 109.00, 96.97, 94.01, 54.93, 54.88.

Method 2(1 step):

To a boiling mixture of *m*-anisidine (1.56 ml, 20.0 mmol) and *N,N*-dimethylaniline (3.5 ml) was added 2-bromo-4-methoxyacetophenone (1.37 g in EtOAc, 6.00 mmol) slowly by syringe. After addition, the mixture was kept at 170° C for 1 hour. The reaction mixture was cooled to room temperature and a dark colored solid was formed. EtOAc was added along with HCl (2 N). The aqueous layer was extracted with EtOAc several times. The combined organic layers were washed with brine, and dried over MgSO_4 . Solvent was removed under the reduced pressure to afford a dark brown colored solid. Purification by recrystallization in EtOH afforded indole 31 as a white crystalline material.

^1H NMR(CDCl_3): δ 11.24 (1H, br, s), 7.72 (2H, d, J 8.82 Hz), 7.36 (1H, d, J 8.57 Hz), 7.00 (2H, d, J = 8.84 Hz), 6.85 (1H, d, J = 2.07 Hz), 6.66 (1H, d, J = 1.66 Hz), 6.63 (1H, dd, J 8.59, 2.28 Hz), 3.78 (3H, s), 3.77 (3H, s).

^{13}C NMR (CDCl_3): δ 158.15, 155.22, 137.44, 136.33, 125.60, 124.93, 122.82, 120.04, 114.07, 109.00, 96.97, 94.01, 54.93, 54.88.

Melting Point: 208-229.5° C

HRMS (EI) M^+ calcd for $\text{CH}_{16}\text{NO}_2$ 253.3035, found 253.1060.

Preparation of Trimethoxybenzoate 2-Phenylindole 33

To a well stirred solution of indole 31 (0.502 g, 1.98 mmol) in *o*-dichlorobenzene (10 ml) was added trimethoxybenzoylchloride (0.692 g, 3.00 mmol). The reaction mixture was heated to reflux for 12 hours. Solvent was removed by distillation under reduced pressure. After cooling down to room temperature, a dark solid formed which was dissolved in chloroform and purified by silica gel column chromatography with chloroform as the eluent. The collected mixture was again purified by column chromatography (50:50 hexanes:EtOAc) affording trimethoxybenzyl

indole 33 (0.744 g, 1.66 mmol, 84%) as a yellow oily gel. Pale yellow-green crystals were obtained by recrystallization from a mixture of ethanol and hexanes.

¹H NMR (CDCl₃): δ 8.63 (1H, br, s), 7.88 (1H, d, *J* = 9.39 Hz), 7.24 (2H, d, *J* = 8.78 Hz), 6.95 (2H, s), 6.90 (2H, m), 6.71 (2H, d, *J* = 8.79 Hz), 3.86 (3H, s), 3.80 (3H, s), 3.73 (3H, s), 3.68 (6H, s);

¹³C NMR (CDCl₃): δ 192.23, 159.73, 157.06, 152.42, 142.85, 141.01, 136.41, 134.65, 130.16, 124.28, 122.94, 122.17, 113.67, 112.46, 111.52, 107.24, 94.54, 60.78, 55.92, 55.54, 55.14.

Melting Point: 153-155°C

Anal. Calcd for C₂₆H₂₅N₆: C, 69.79; H, 5.63; N, 3.13. Found: C, 69.61; H, 5.63; N, 3.01.

EXAMPLE 2

INHIBITION OF TUBULIN POLYMERIZATION ASSAY

IC₅₀ values for tubulin polymerization were determined according to the procedure described in Bai et al. Purified tubulin is obtained from bovine brain cells as described in Hamel and Lin. Various amounts of inhibitor were preincubated for 15 minutes at 37° C with purified tubulin. After the incubation period, the reaction was cooled and GTP was added to induce tubulin polymerization. Polymerization was then monitored in a Gilford spectrophotometer at 350 nm. The final reaction mixtures (0.25 ml) contained 1.5 mg/ml tubulin, 0.6 mg/ml microtubule-associated proteins (MAPs), 0.5 mM GTP, 0.5 mM MgCl₂, 4% DMSO and 0.1M 4-morpholineethanesulfonate buffer (MES, pH 6.4). IC₅₀ is the amount of inhibitor needed to inhibit tubulin polymerization 50% with respect to the amount of inhibition that occurs in the absence of inhibitor. The IC₅₀ value determined for 3-(3',4',5'-trimethoxybenzoyl)-2-(4'-methoxyphenyl)-6-methoxyindole was 0.5-1.5 μM.

EXAMPLE 3

CYTOTOXIC ASSAY WITH P388 LEUKEMIA CELLS

One of the newly prepared compounds was evaluated for cytotoxic activity against P388 leukemia cells using an assay system similar to the National Cancer Institute procedure described below and in Monks et al. The ED₅₀ value (defined as the effective dosage required to inhibit 50% of cell growth) of 3-(3',4',5' trimethoxybenzoyl)-2-(4'-methoxyphenyl)-6-methoxyindole was found to be 0.0133 µg/mL.

EXAMPLE 4

GROWTH INHIBITORY ACTIVITY AGAINST OTHER
CANCER CELL LINES

3-(3',4',5'-Trimethoxybenzoyl)-2-(4'-methoxyphenyl)-6-methoxyindole was evaluated in terms of growth inhibitory activity against several human cancer cell lines, including pancreas, ovarian, CNS, lung-NSC, colon, and prostate lines. The assay used is described in Monks et al. Briefly, the cell suspensions, diluted according to the particular cell type and the expected target cell density (5,000-40,000 cells per well based on cell growth characteristics), were added by pipet (100 µl) to 96-well microtiter plates. Inoculates were allowed a preincubation time of 24-28 hours at 37°C for stabilization. Incubation with the inhibitor compounds lasted for 48 hours in 5% CO₂ atmosphere and 100% humidity. Determination of cell growth was done by in situ fixation of cells, followed by staining with a protein-binding dye, sulforhodamine B (SRB), which binds to the basic amino acids of cellular macromolecules. The solubilized stain was measured spectrophotometrically. The results of these assays are shown in Table 1. GI₅₀ is defined as the dosage required to inhibit tumor cell growth by 50%.

Table IV. Activity of Indole Ligand Against Selected Human Cancer Cell lines (*In Vitro*).

Indole-based Ligand 33		
CELL TYPE	CELL LINE	GI ₅₀ (μG/mL)
Pancreas-a	BXPC-3	2.0 x 10 ⁻³
Ovarian	OVCAR-3	2.4 x 10 ⁻³
CNS	SF-295	2.4 x 10 ⁻³
Lung-NSC	NCI-H460	2.6 x 10 ⁻³
Colon	KM20L2	1.7 x 10 ⁻³
Prostate	DU-145	2.3 x 10 ⁻³

Indole and indole containing compounds of therapeutic efficacy have been known for many, many years. What is truly unique about the indole compounds described in this application is the fact that these compounds are the first (to the best of our knowledge) indole-based ligands to incorporate the 3,4,5-trimethoxyaryl motif reminiscent of colchicine and combretastatin A-4 arranged in an appropriate molecular conformation such that a pseudo aryl-aryl pi stacking interaction can take place. It is our contention that such an aryl-aryl interaction of the appropriate centroid-to-centroid distance (approximately 4.7 Å) is imperative for enhanced binding affinity to the colchicine site on β-tubulin. It is this binding that ultimately leads to an inhibition of tubulin polymerization which manifests itself as a cytotoxic event. It should be readily apparent to any practitioner skilled in the art that there are various ways of appending trimethoxyaryl and trimethoxyaroyl groups around an indole molecular scaffold in a manner which will result in a similar molecular conformation capable of undergoing pseudo pi-pi stacking. In addition, although the trimethoxyaryl motif seems optimal for enhanced tubulin binding, it is also very possible that another combination of alkoxy substituents (such as ethoxy, propoxy, isopropoxy, allyloxy, etc.) either as a trisubstituted pattern or as disubstituted (with one type of alkoxy moiety) and monosubstituted (with a different alkoxy moiety), or with three distinct types of alkoxy moieties may also have good tubulin binding characteristics. It is also conceivable that instead of having aryl alkoxy groups, it may be possible to substitute simply aryl-alkyl and aryl-alkenyl moieties and still maintain the enhanced cytotoxicity profile. Phenolic groups may also have activity on these described indole ligands. The synthesis of any of these modified

indole-ligands will be very straight-forward for anyone skilled in the art, and often will only involve a different choice of initial starting materials. To prepare these alternative ligands, the same synthetic schemes (Figures 6, 9, 11, 12-16), or similar schemes with only slight modifications may be employed. In previous studies with the benzo[*b*]thiophene ligands, we
5 have demonstrated that the carbonyl group can be replaced with an oxygen to generate a new compound which maintains the same or similar biological efficacy with tubulin. Similarly, the replacement of the carbonyl group in the described indole ligand may be replaced with an oxygen atom (ether linkage) to generate a new derivative which would be predicted to have good activity with tubulin. This compound may be prepared by an addition elimination reaction utilizing the
10 trimethoxyphenolic anion as a nucleophile as described by us for the benzo[*b*]thiophene compounds. Other linkage atoms between the aryl aryl rings are conceivable as well.

All of the compositions and methods disclosed and claimed herein can be made and executed without undue experimentation in light of the present disclosure. While the compositions and methods of this invention have been described in terms of preferred
15 embodiments, it will be apparent to those of skill in the art that variations may be applied to the compositions and/or methods and in the steps or in the sequence of steps of the method described herein without departing from the concept, spirit and scope of the invention. More specifically, it will be apparent that certain agents which are both chemically and physiologically related may be substituted for the agents described herein while the same or similar results would be
20 achieved. All such similar substitutes and modifications apparent to those skilled in the art are deemed to be within the spirit, scope and concept of the invention as defined by the appended claims.

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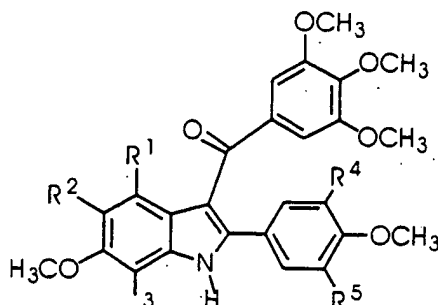
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What is claimed is:

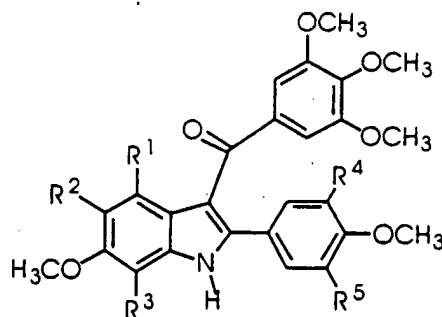
1. A compound of the structure:



wherein

R^1 through R^5 contain at least one phenolic moiety or at least one amine group (NH_2 , NHR^1 , or NR^6R^7 where R^6 and R^7 are the same or different alkyl having up to 8 carbon atoms), benzyl, or aryl while the remaining R^1 through R^5 are hydrogen.

2. A compound of the structure:

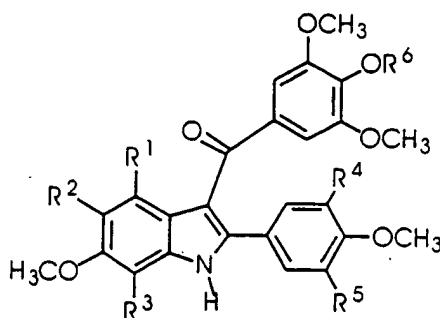


wherein

R^1 through R^5 contain at least one phosphate ester moiety ($-OP(O)(OM^+)_2$) or a phosphoramidate ($-NP(O)(OM^+)_2$) where M is a cation or ($-NP(O)(OR)_2$) where

R is an alkyl with up to 8 carbon atoms (the two R groups are the same or different, benzyl, or aryl while the remaining R¹ through R⁵ are hydrogen.

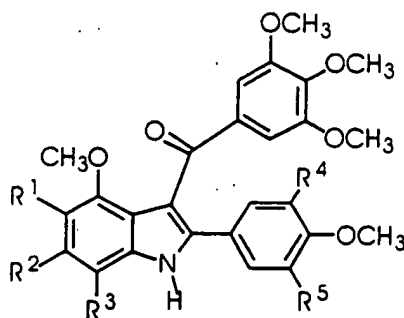
3. A compound of the structure:



wherein

R¹ through R⁵ contain at least one phosphate ester moiety (-OP(O)(O⁻M⁺)₂) or a phosphoramidate (-NP(O)(O⁻M⁺)₂) where M is a cation or (-NP(O)(OR)₂) where R is an alkyl with up to 8 carbon atoms (the two R groups are the same or different), benzyl, or aryl while the remaining R¹ through R⁵ are hydrogen, and R⁶ is hydrogen or alkyl.

4. A compound of the structure:

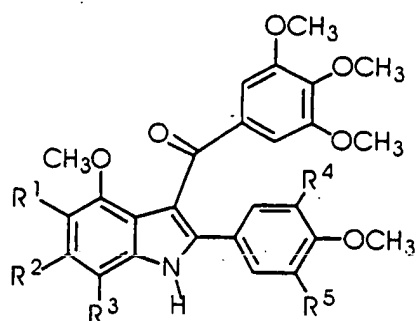


wherein

R¹ through R⁵ contain at least one phenolic moiety or at least one amine (NH₂, NHR¹, or NR⁶R⁷ where R⁶ and R⁷ the same or different alkyl having up to 8

carbon atoms, benzyl, or aryl groups) while the remaining R^1 through R^5 are a hydrogen.

5. A compound of the structure:

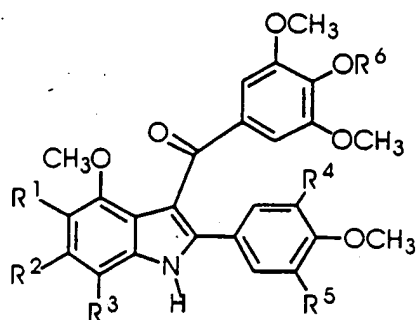


wherein

5

R^1 through R^5 contain at least one phosphate ester moiety ($-\text{OP}(\text{O})(\text{O}^-\text{M}^+)_2$) or a phosphoramidate ($-\text{NP}(\text{O})(\text{O}^-\text{M}^+)_2$) where M is a cation or ($-\text{NP}(\text{O})(\text{OR})_2$) where R is an alkyl with up to 8 carbon atoms (the two R groups are the same or different), benzyl, or aryl while the remaining R^1 through R^5 are hydrogen.

6. A compound of the structure:



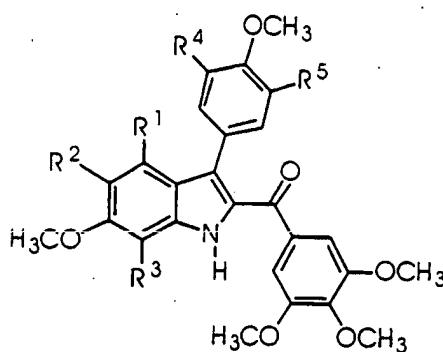
wherein

5

R^1 through R^5 contain at least one phosphate ester moiety ($-\text{OP}(\text{O})(\text{O}^-\text{M}^+)_2$) or a phosphoramidate ($-\text{NP}(\text{O})(\text{O}^-\text{M}^+)_2$) where M = a cation or ($-\text{NP}(\text{O})(\text{OR})_2$) where

R is an alkyl with up to 8 carbon atoms (the two R groups are the same or different), or benzyl, or aryl groups, while the remaining R¹ through R⁵ are a hydrogen and R⁶ is hydrogen or alkyl.

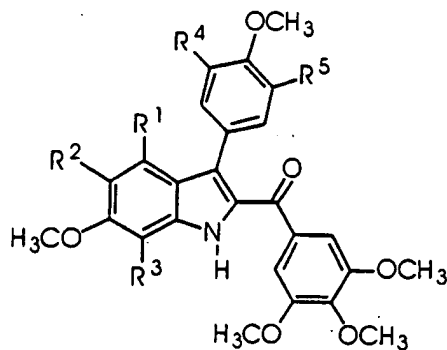
7. A compound of the structure:



wherein

R¹ through R⁵ contain at least one phenolic moiety or at least one amine group (NH₂, NHR or NR⁶R⁷ where R⁶ and R⁷ are the same or different alkyl having up to 8 carbon atoms may be the same or different), or benzyl, or aryl groups) while the remaining R¹ through R⁵ are a hydrogen.

8. A compound of the structure:

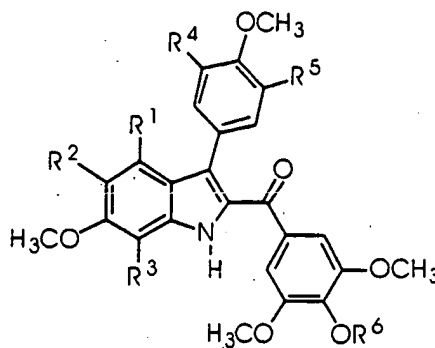


wherein

5

R^1 through R^5 contain at least one phosphate ester $(-OP(O)(O^+M^-)_2)$ or a phosphoramidate $(-NP(O)(O^+M^-)_2)$ where M is a cation or $(-NP(O)(OR)_2)$ where R is an alkyl with up to 8 carbon atoms (the two R groups are the same or different), benzyl, or aryl while the remaining R^1 through R^5 are hydrogen.

9. A compound of the structure:

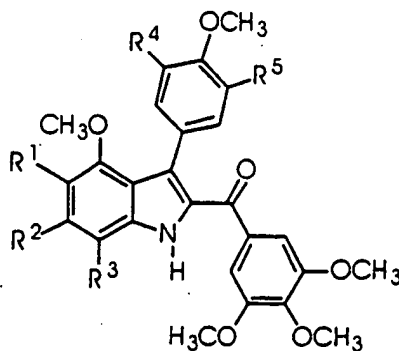


wherein

5

R^1 through R^5 contain at least one phosphate ester $(-OP(O)(O^+M^-)_2)$ or phosphoramidate $(-NP(O)(O^+M^-)_2)$ where M is a cation or $(-NP(O)(OR)_2)$ where R is an alkyl with up to 8 carbon atoms (the two R groups are the same or different), benzyl, or aryl, while the remaining R^1 through R^5 are hydrogen, and R^6 is hydrogen or alkyl.

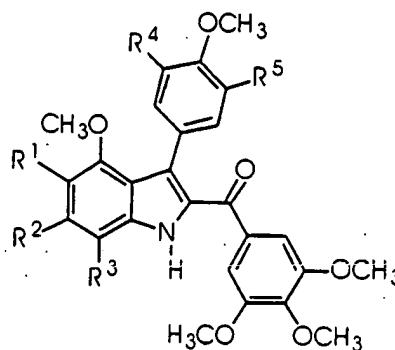
10. A compound of the structure:



wherein

- 5 R^1 through R^5 contain at least one phenolic moiety or at least one amine group (NH_2 , NHR^1 , or NR^6R^7 where R^6 and R^7 are the same or different alkyl having up to 8 carbon atoms, benzyl, or aryl) while the remaining R^1 through R^5 are a hydrogen.

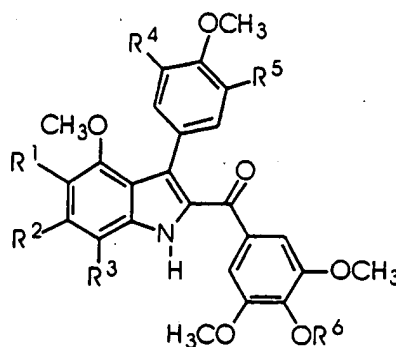
11. A compound of the structure:



wherein

- 5 R^1 through R^5 contain at least one phosphate ester ($-OP(O)(O^+M^-)_2$) or a phosphoramidate ($-NP(O)(O^+M^-)_2$) where M is a cation or ($-NP(O)(OR)_2$) where R is an alkyl with up to 8 carbon atoms (the two R groups are the same or different), benzyl, or aryl, while the remaining R^1 through R^5 are hydrogen.

12. A compound of the structure:

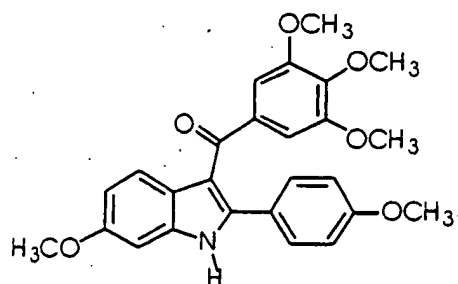


wherein

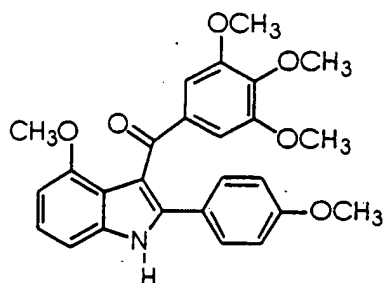
- 5 R^1 through R^5 contain at least one phosphate ester moiety ($-OP(O)(O^+M^-)_2$) or a phosphoramidate ($-NP(O)(O^+M^-)_2$) where M is a cation or ($-NP(O)(OR)_2$) where R is an

alkyl with up to 8 carbon atoms (the two R groups are the same or different), benzyl, or aryl while the remaining R¹ through R⁵ are hydrogen, and R⁶ is hydrogen or alkyl.

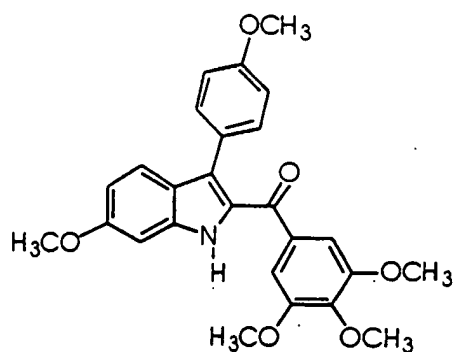
13. A compound of the structure:



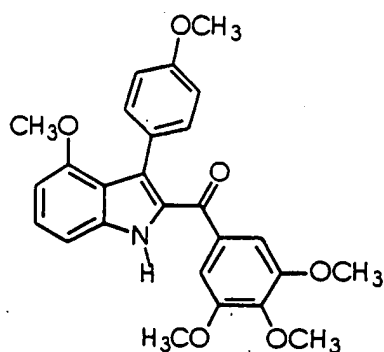
14. A compound of the structure:



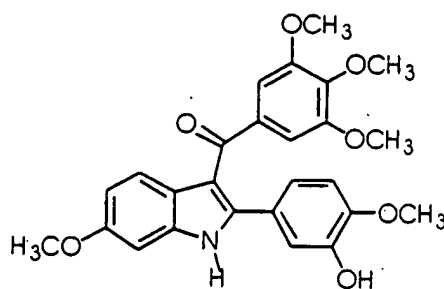
15. A compound of the structure:



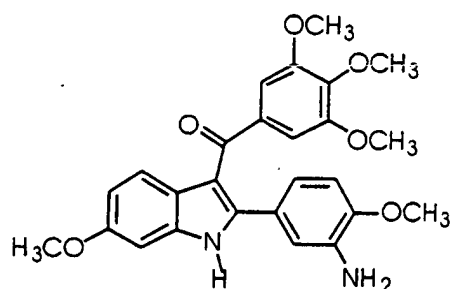
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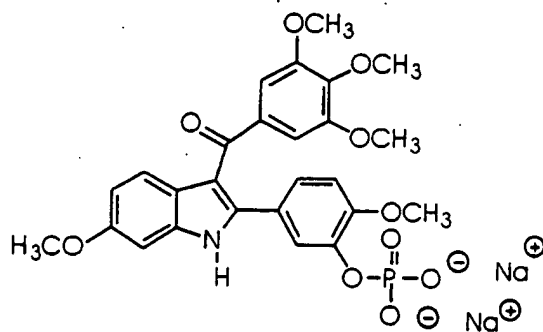
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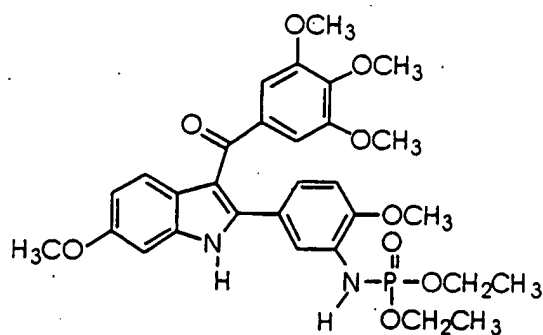
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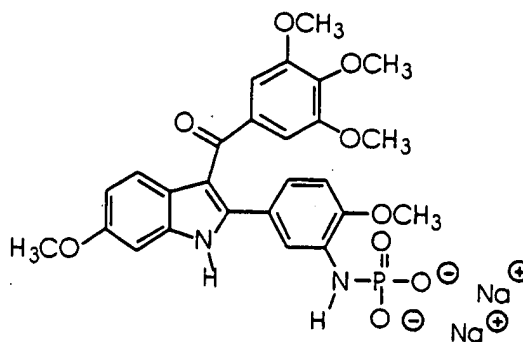
19. A compound of the structure:



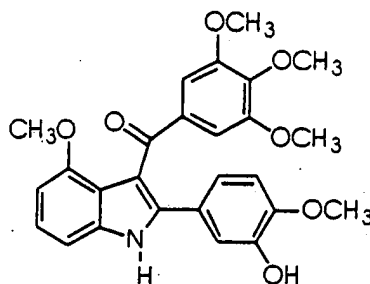
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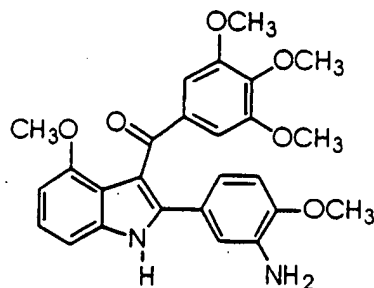
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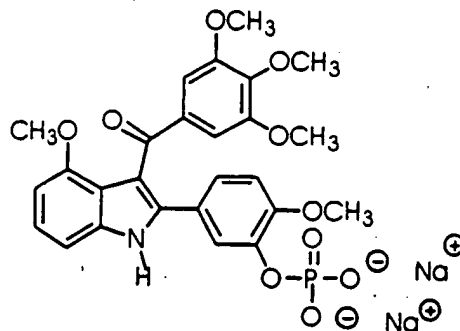
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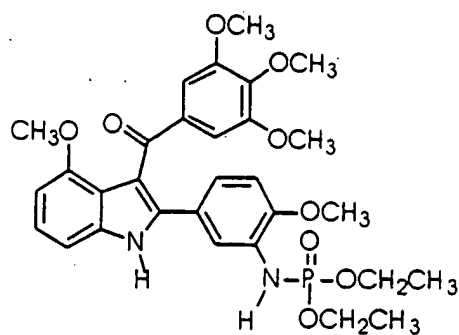
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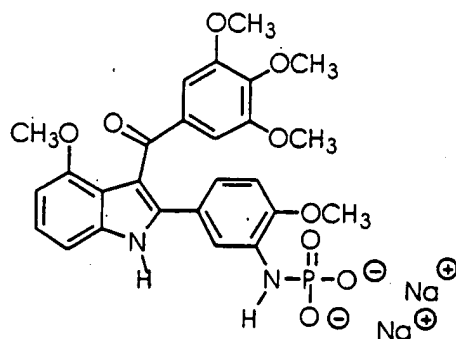
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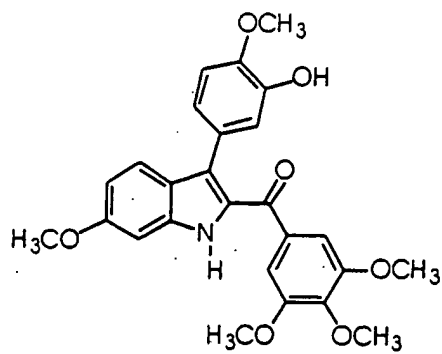
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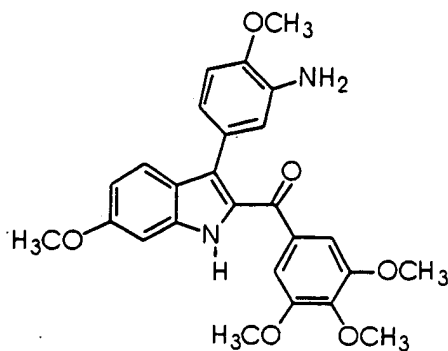
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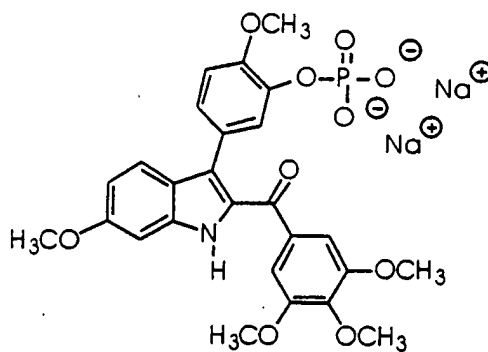
27. A compound of the structure:



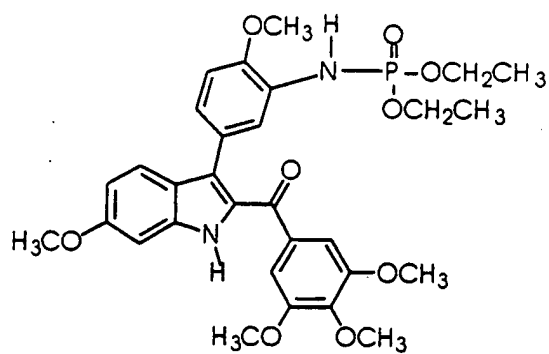
28. A compound of the structure:



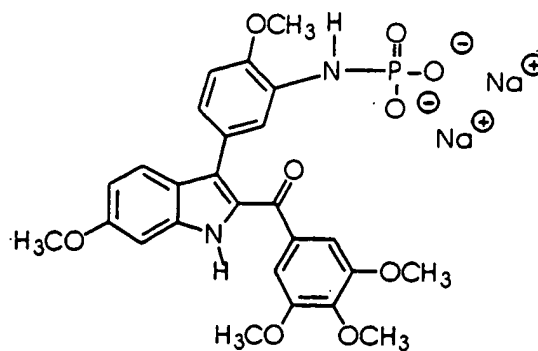
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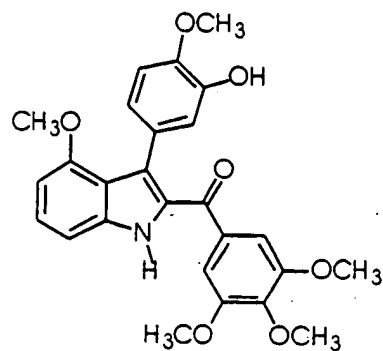
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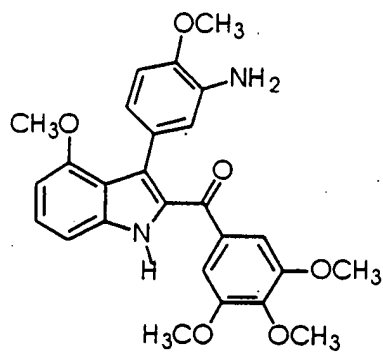
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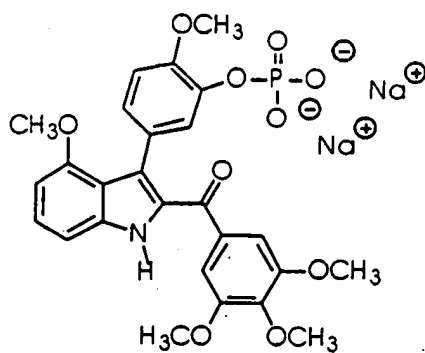
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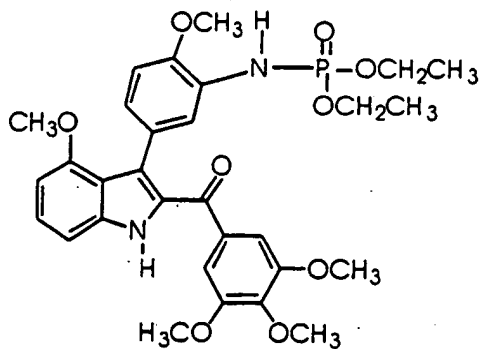
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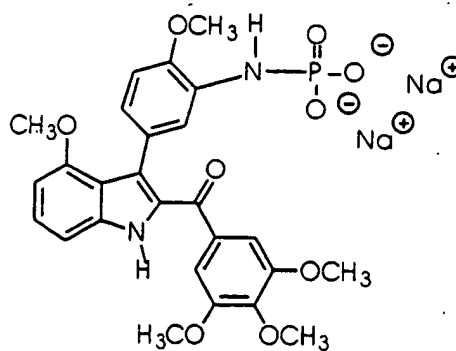
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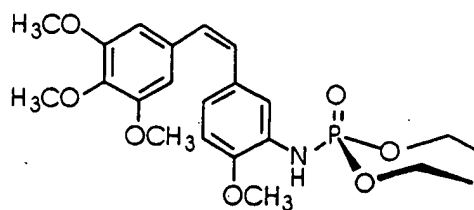
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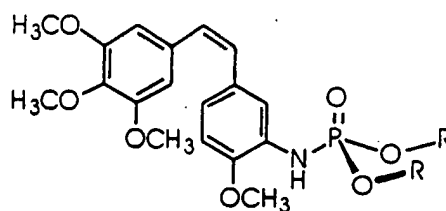
36. A compound of the structure:



37. A compound of the structure:



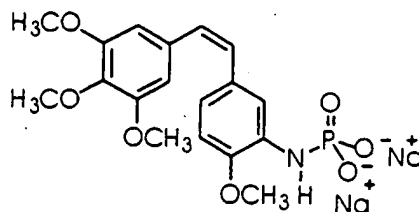
38. A compound of the structure:



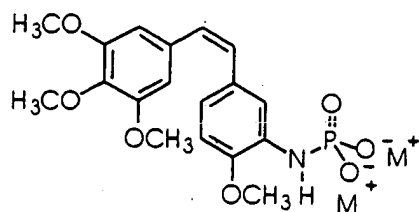
wherein

R is chosen to be any appropriate alkyl or branched alkyl having up to 8 carbon atoms, the two R groups may be the same or different.

39. A compound of the structure:



40. A compound of the structure:



wherein

M^+ is a cation.

41. A method for inhibiting tubulin polymerization by contacting a tubulin-containing system with an effective amount of a compound described in any of claims 1-40.
42. The method of claim 41 wherein said system is in a tumor cell.
43. A method of treating a host afflicted with a neoplastic disease by administering to said host a compound described in any of claims 1-40.
44. The method of claims 41, wherein the contacted system is located in a patient.
45. The method of claim 41 described further as for treating cancer, wherein said cancer may be chosen from the group containing leukemia, lung, colon, thyroid, CNS, melanoma, ovarian, renal, prostate, and breast cancers.

46. A preparation for pharmaceutical use containing a compound from any of claims 1-40 as an active component along with a pharmaceutically acceptable carrier.

47. A method for selectively targeting and destroying tumor vasculature comprising administering an effective amount of a compound described in any of claims 1-40.

FIG. 1

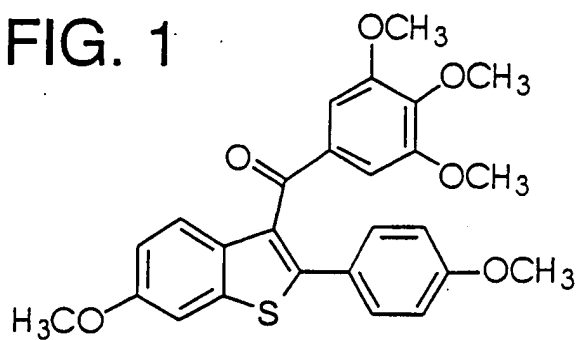


FIG. 2

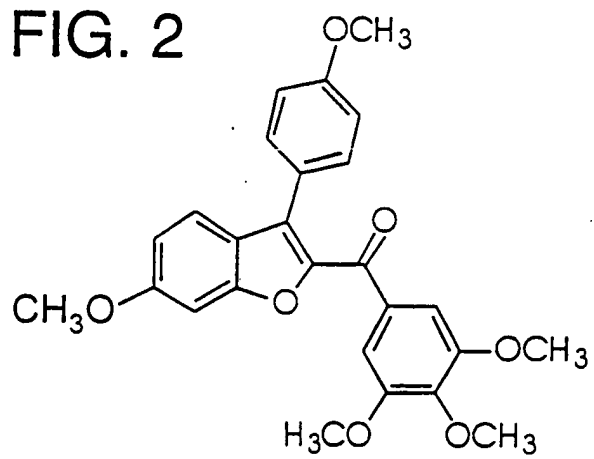


FIG. 3

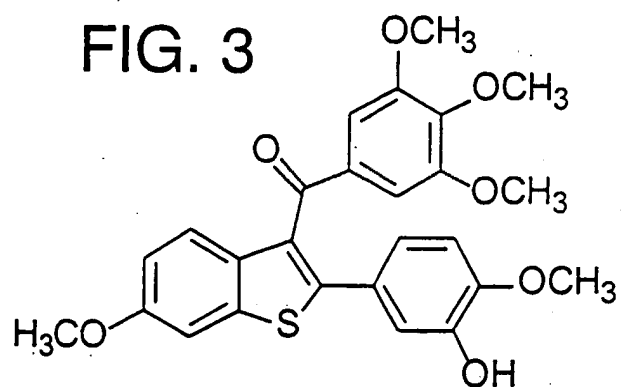


FIG. 4

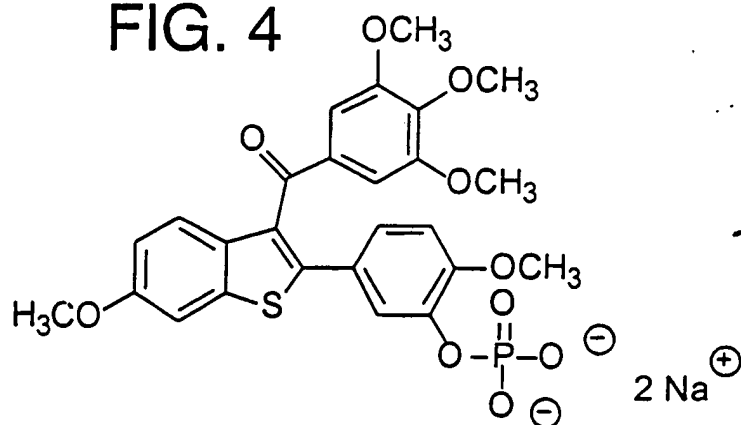


FIG. 5

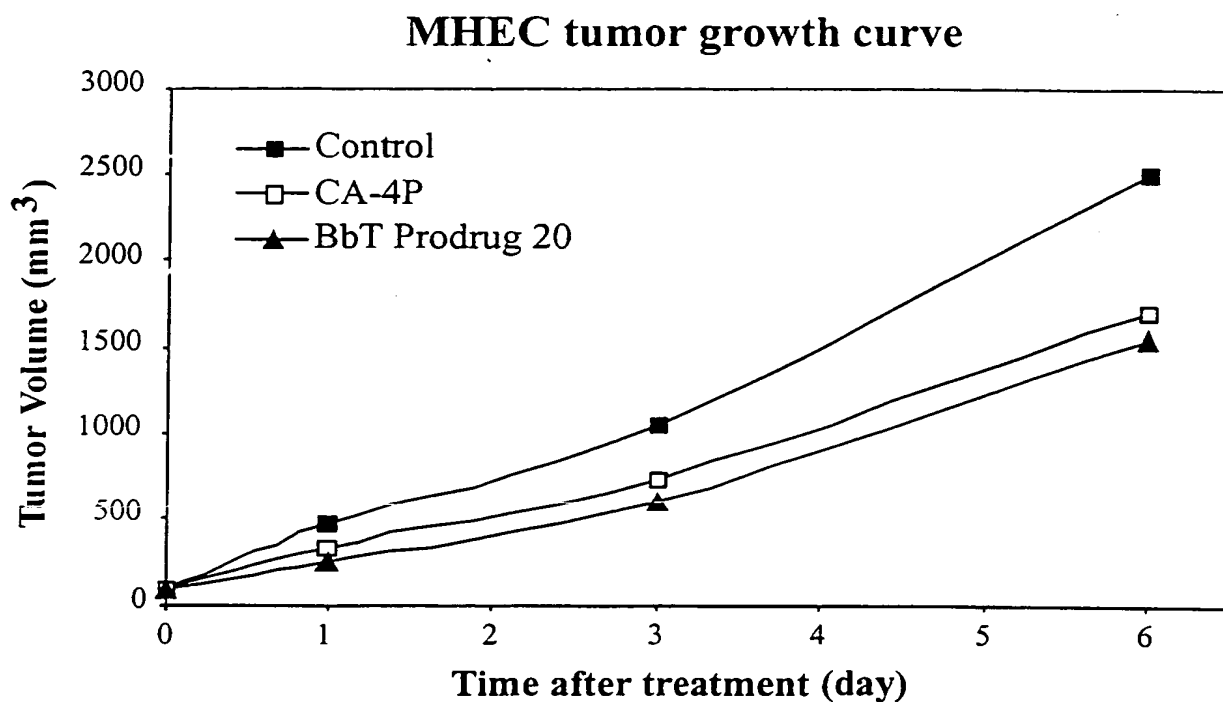


FIG. 6

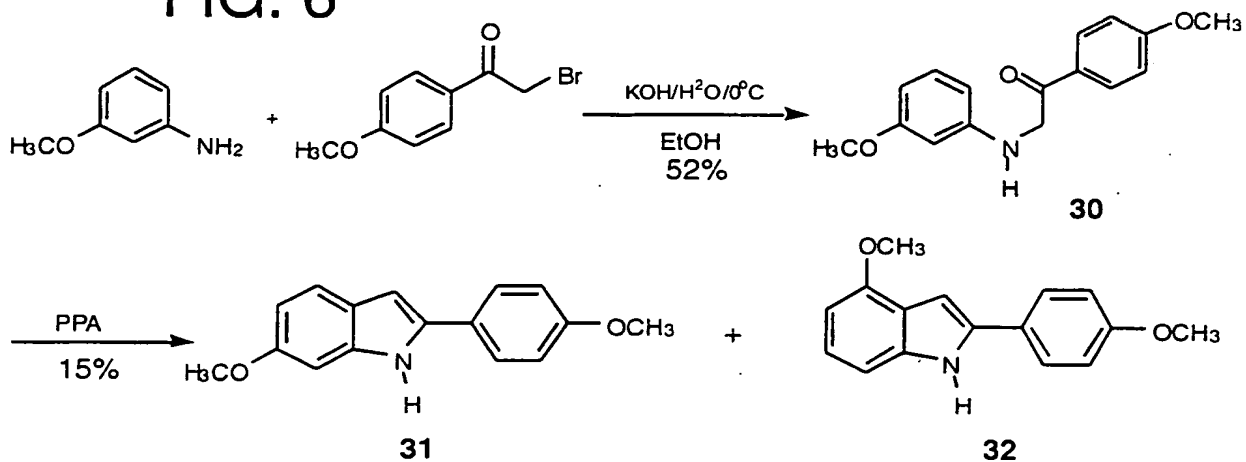


FIG. 7

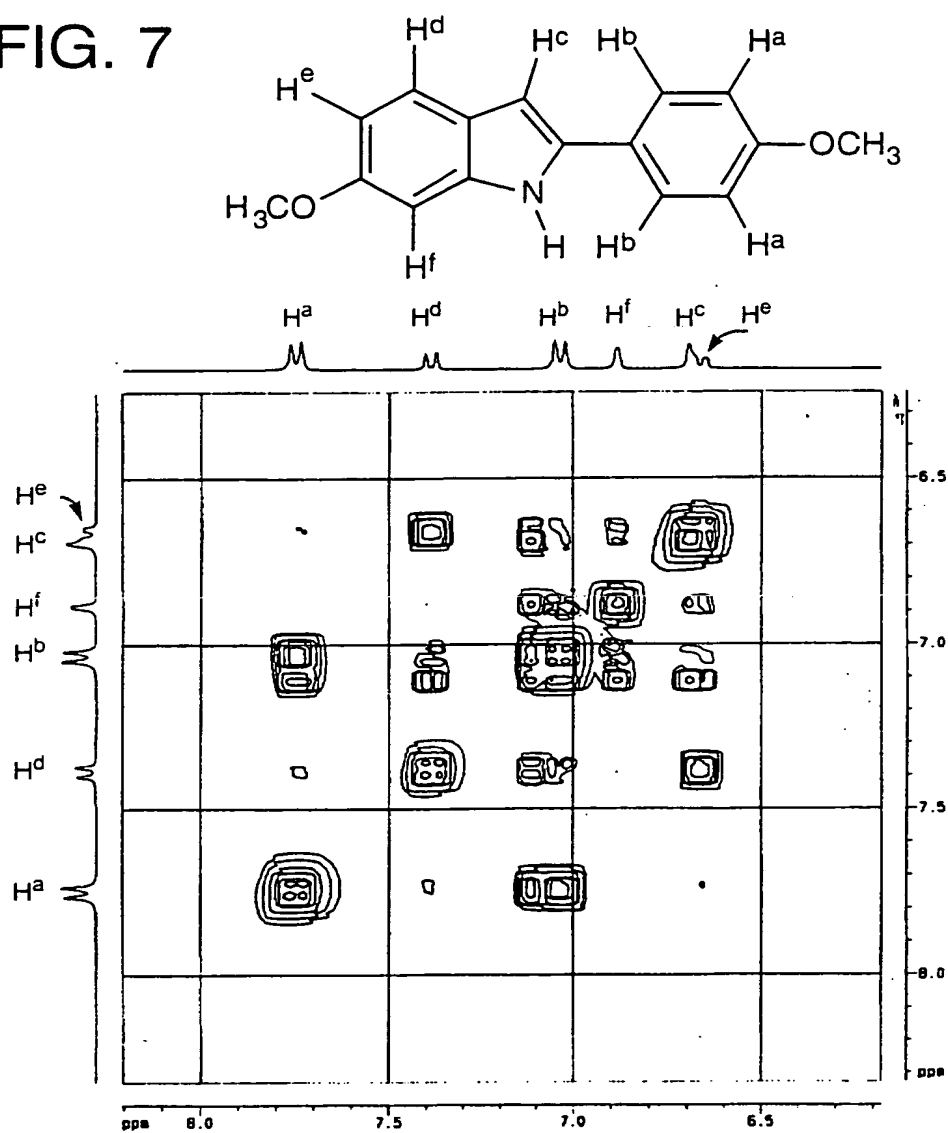


FIG. 8

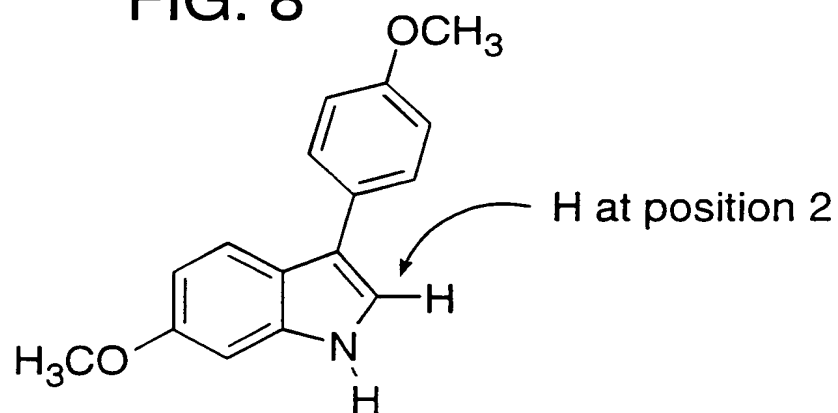


FIG. 9

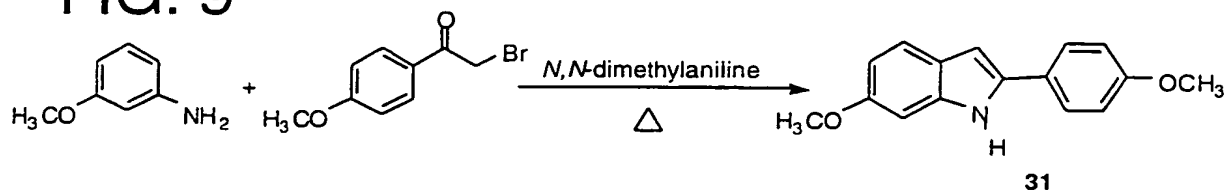


FIG. 10

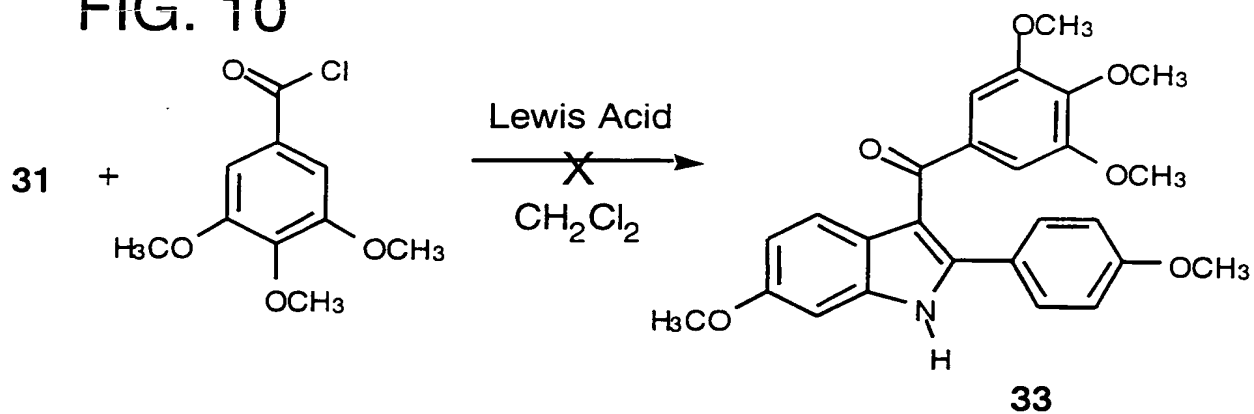


FIG. 11

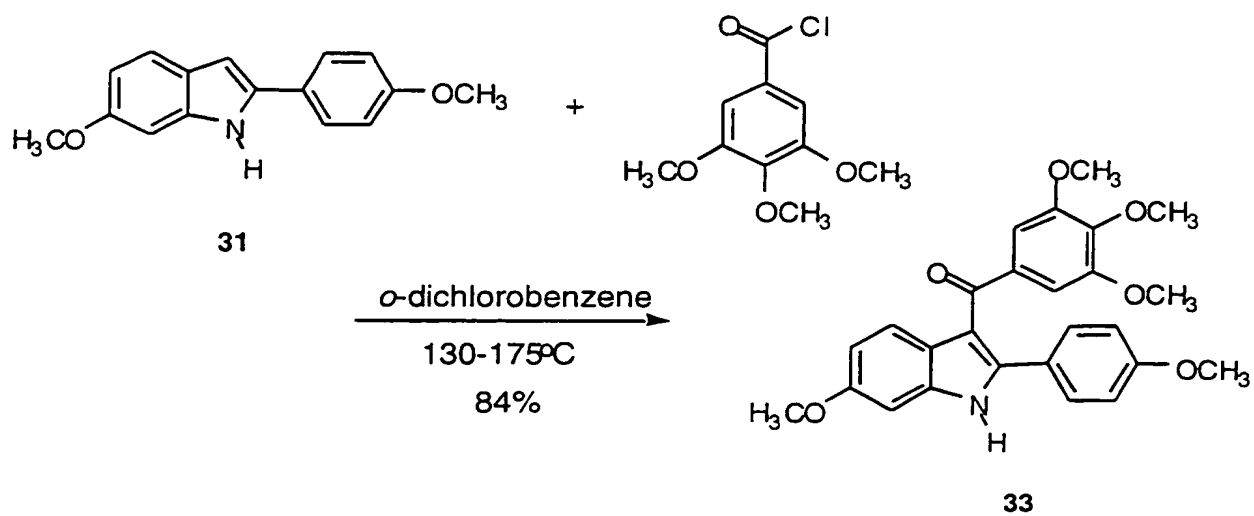


FIG. 12

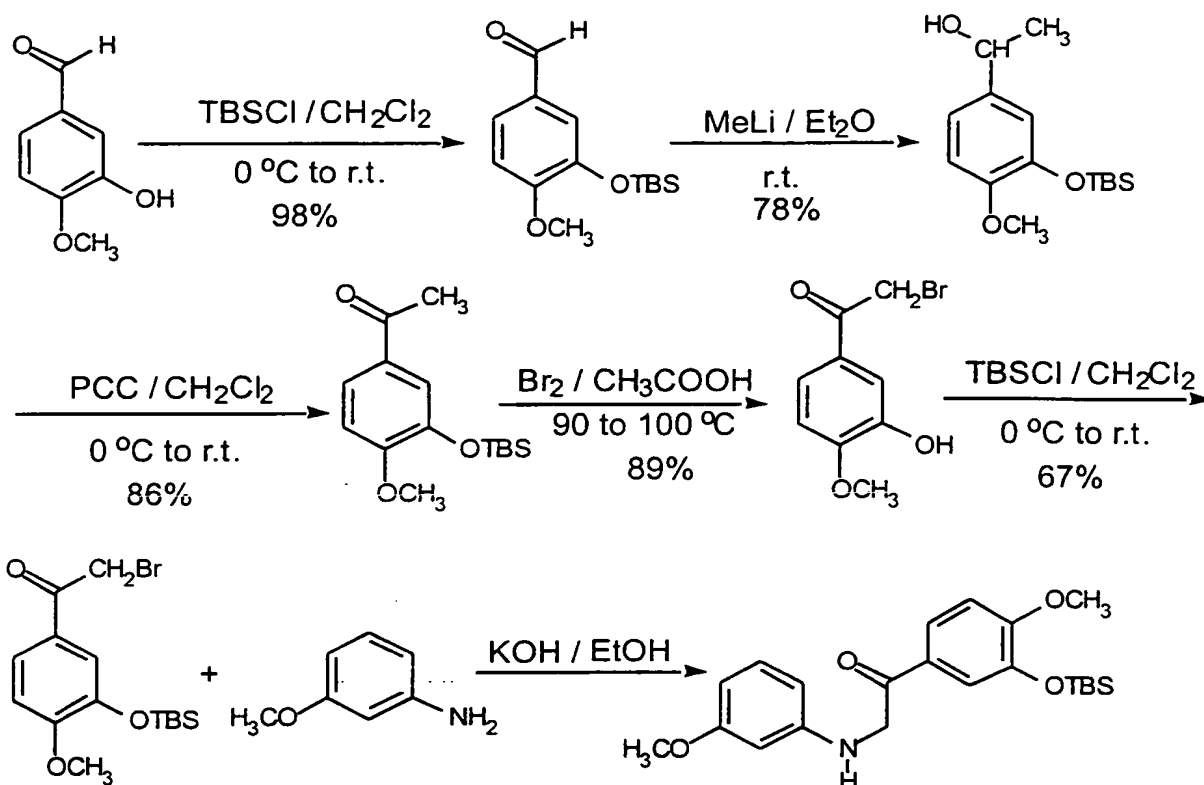


FIG. 13

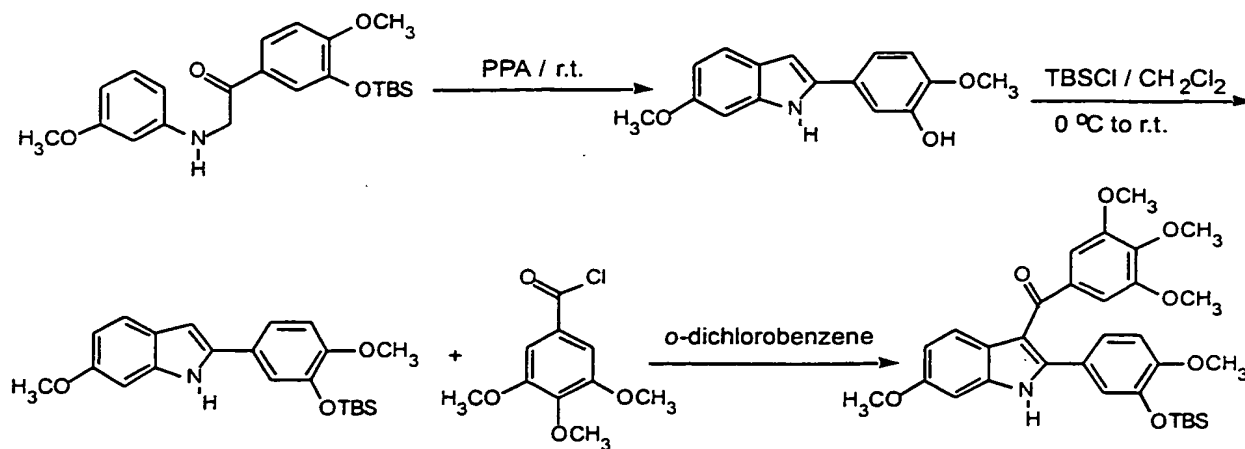


FIG. 14

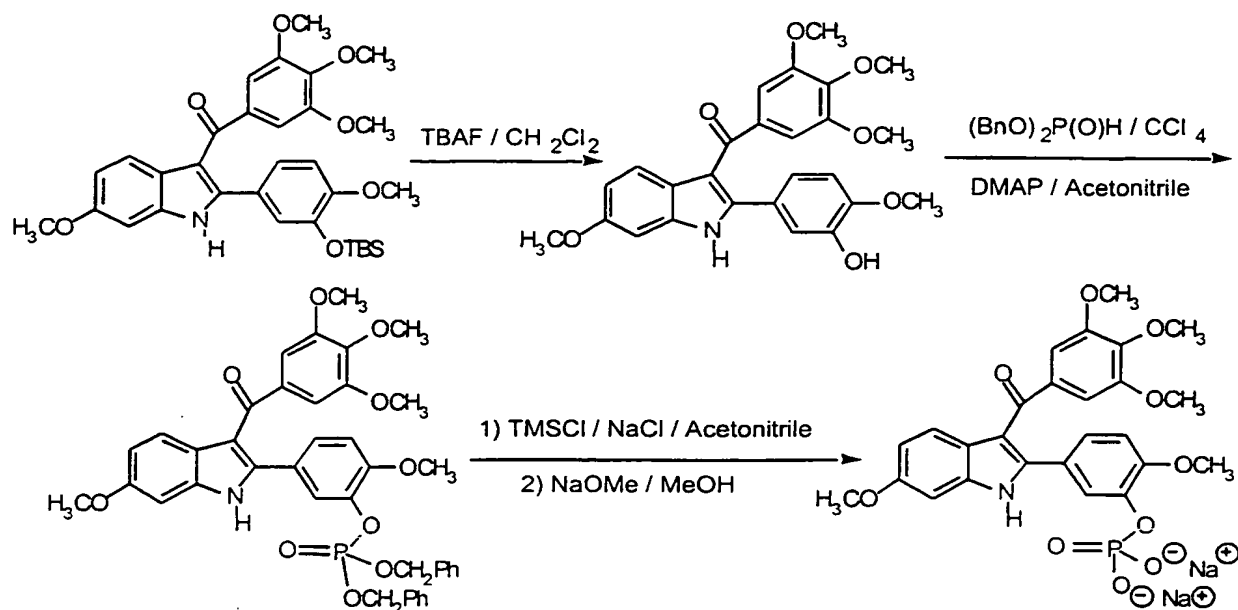


FIG. 15

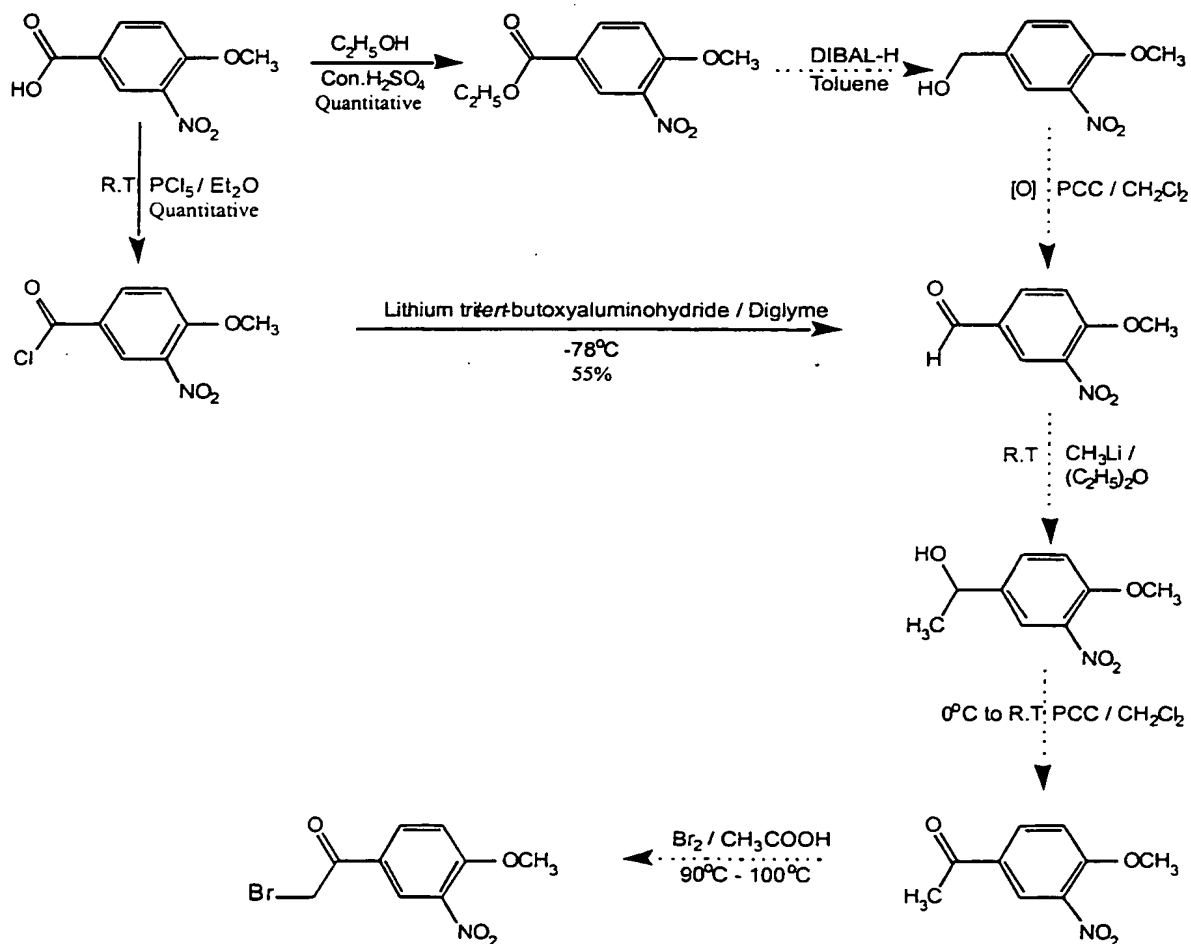


FIG. 16

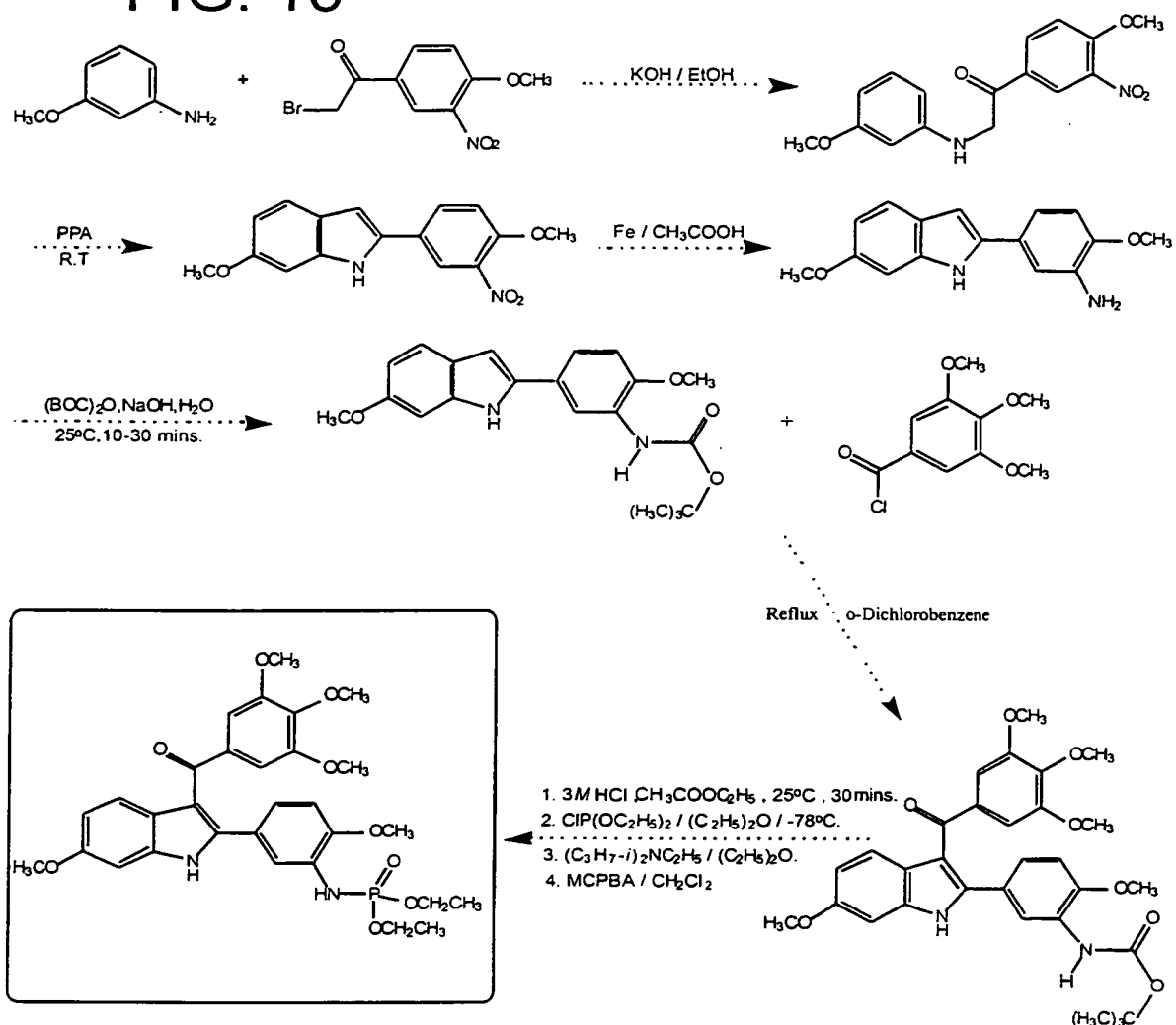


FIG. 17A

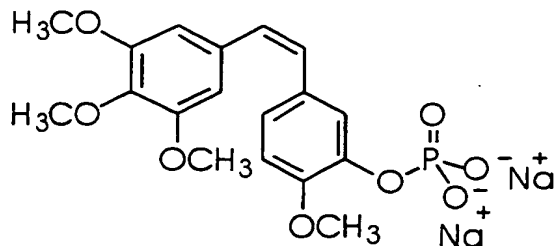


FIG. 17B

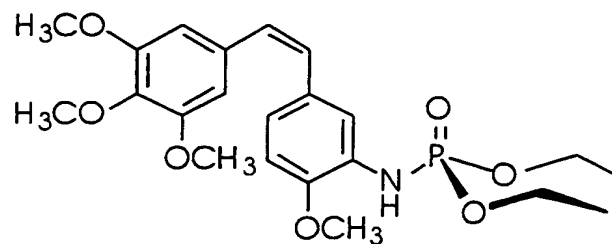


FIG. 18

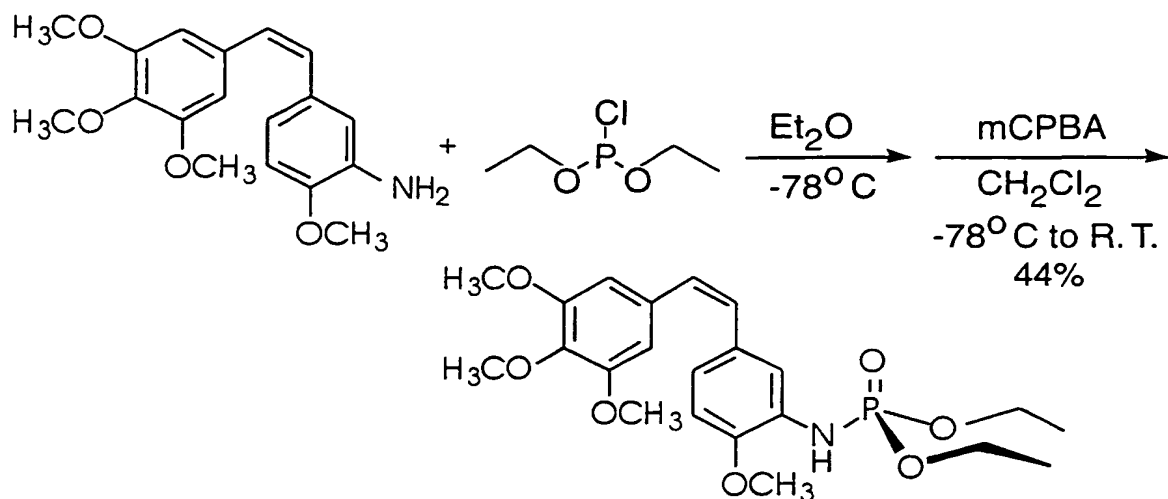


FIG. 19

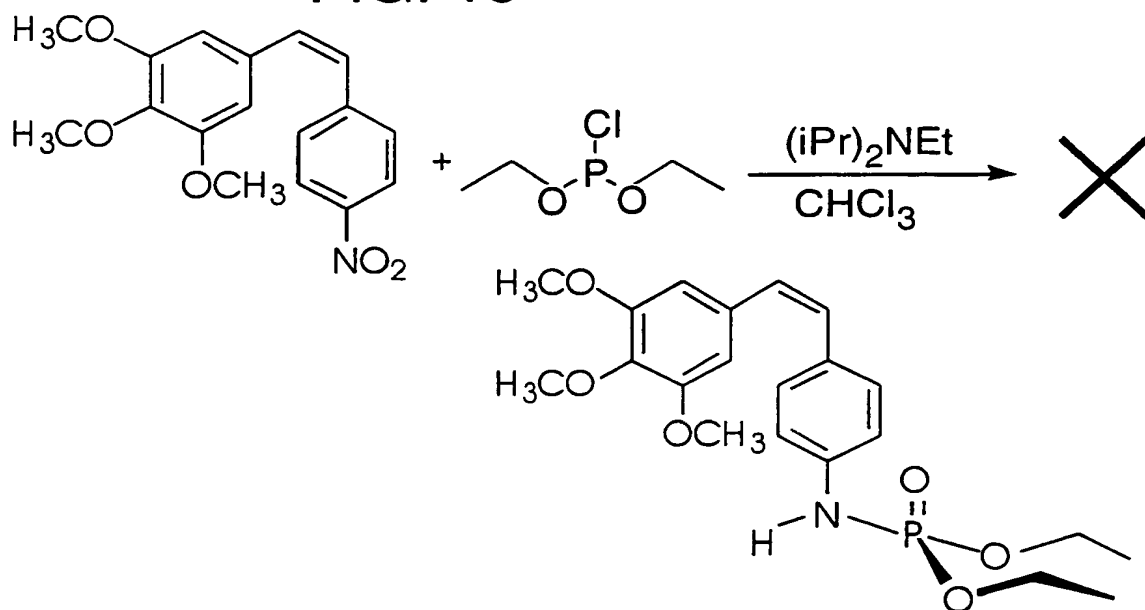


FIG. 20

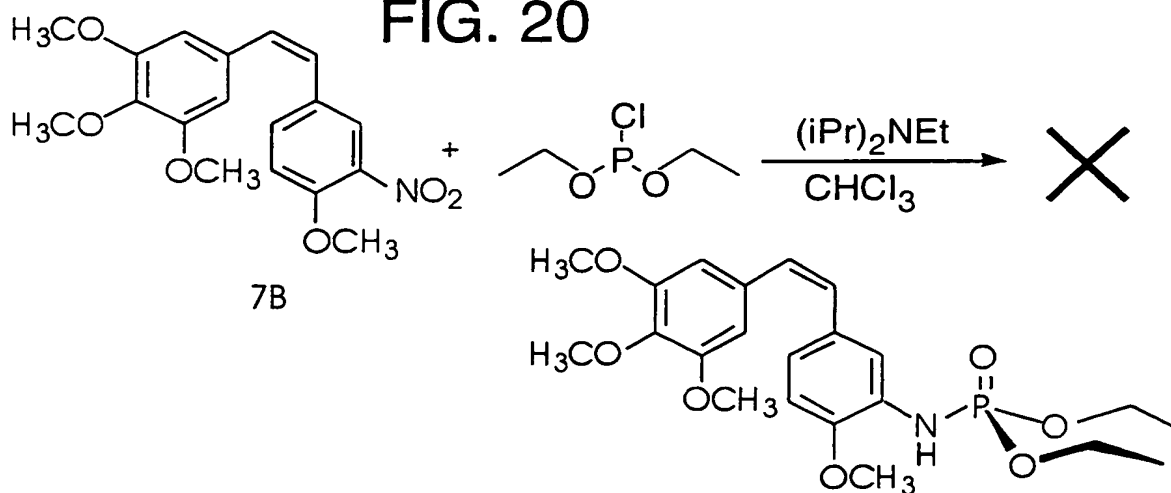


FIG. 21

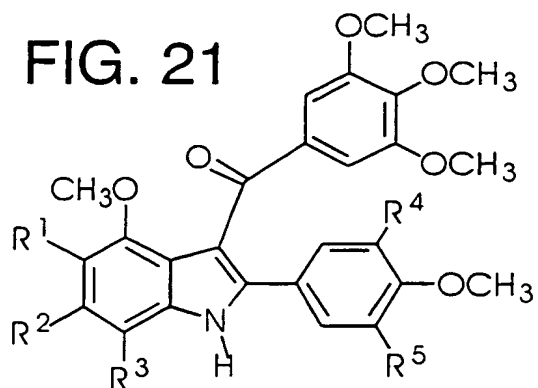


FIG. 22

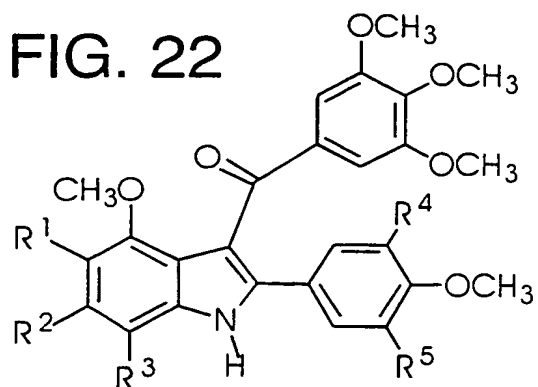


FIG. 23

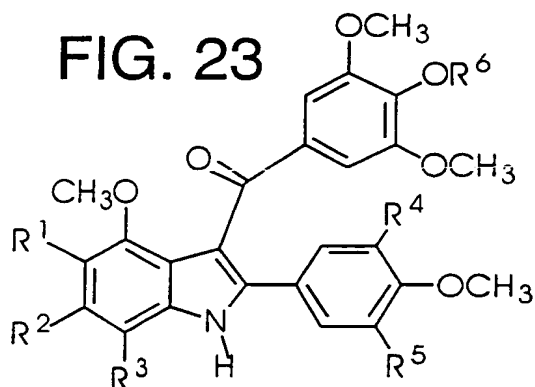


FIG. 24

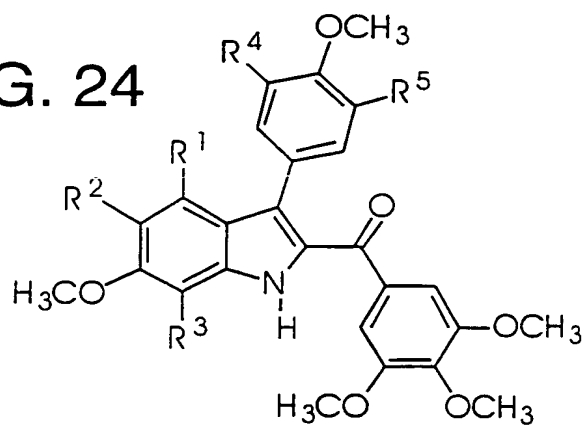


FIG. 25

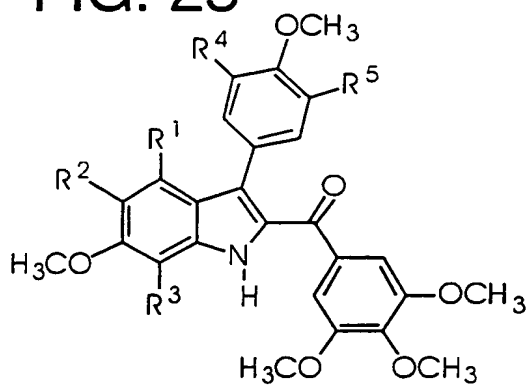


FIG. 26

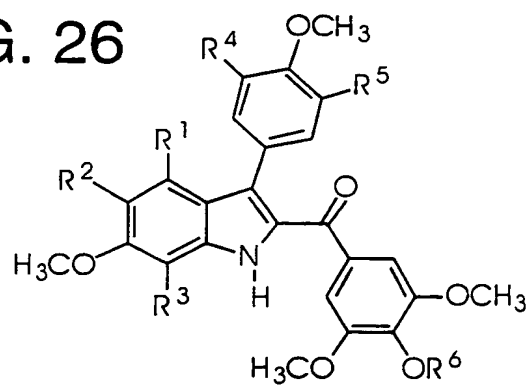


FIG. 27

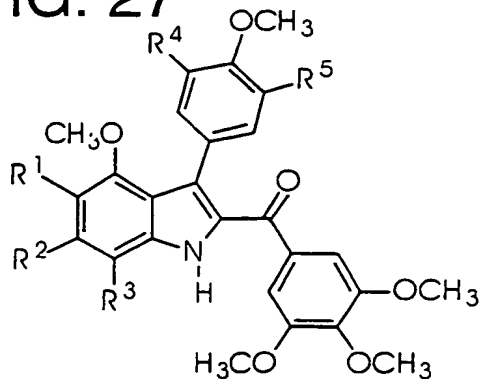


FIG. 28

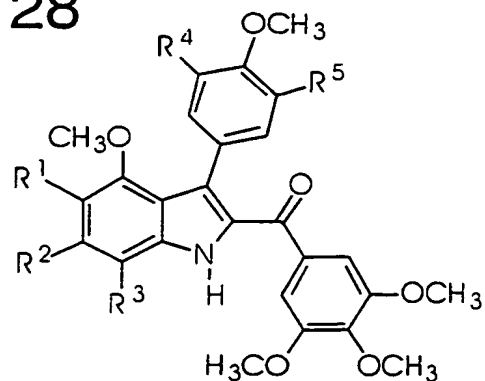


FIG. 29

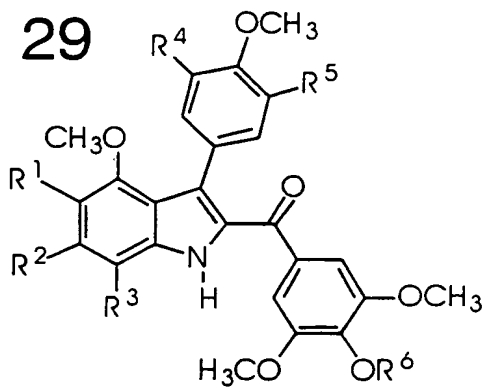


FIG. 30

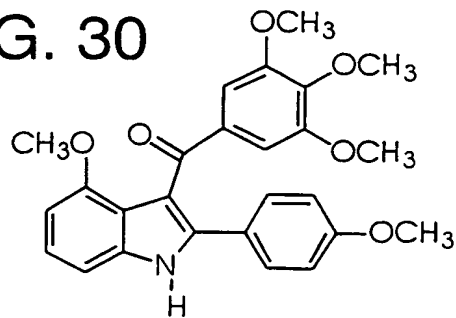


FIG. 31

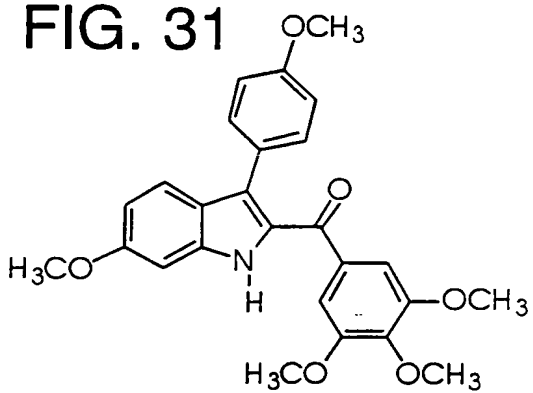


FIG. 32

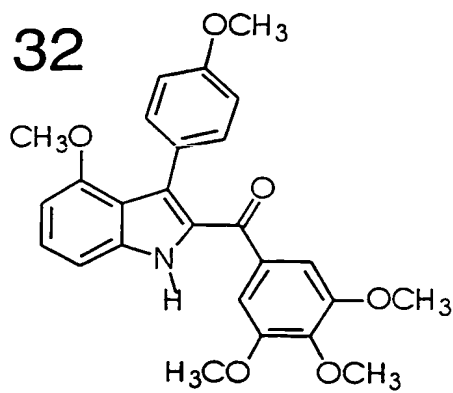


FIG. 33

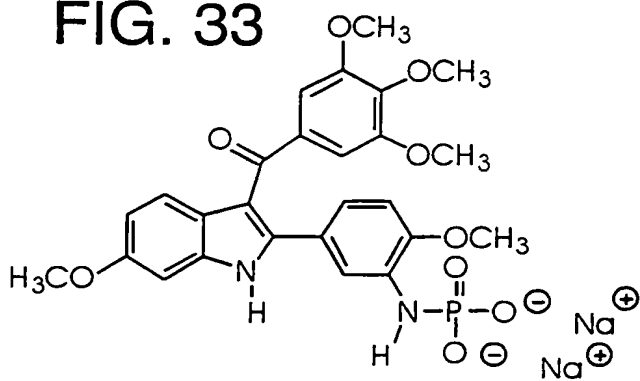


FIG. 34

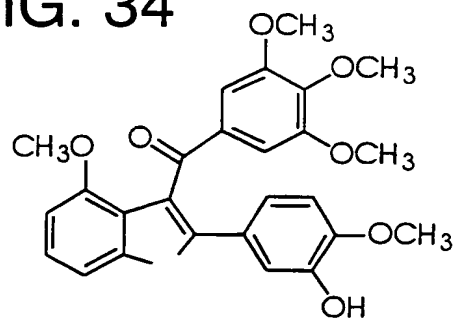


FIG. 35

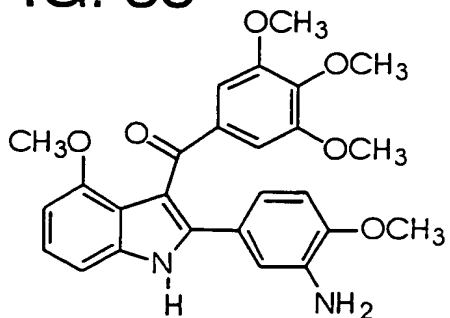


FIG. 36

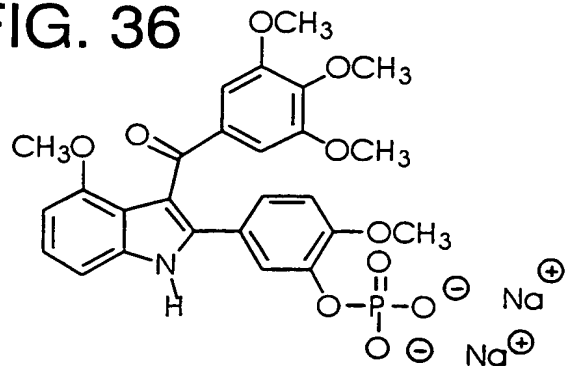


FIG. 37

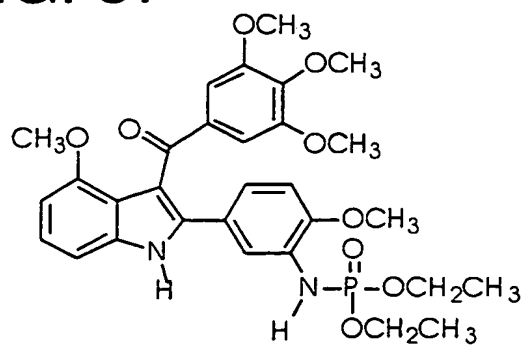


FIG. 38

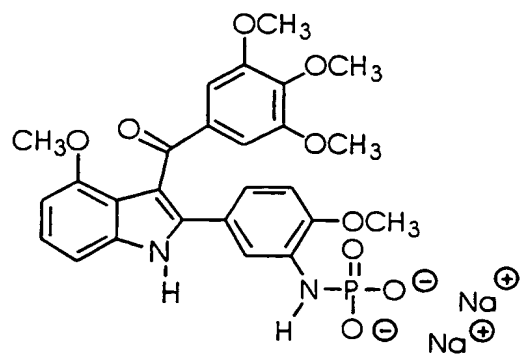


FIG. 39

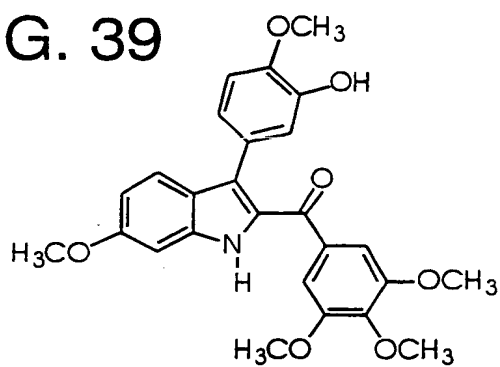


FIG. 40

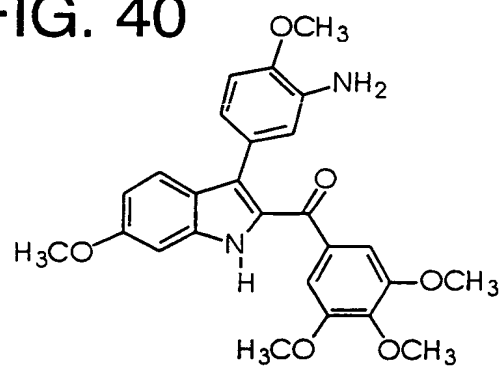


FIG. 41

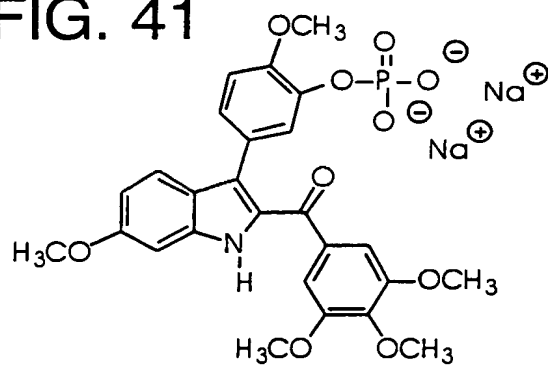


FIG. 42

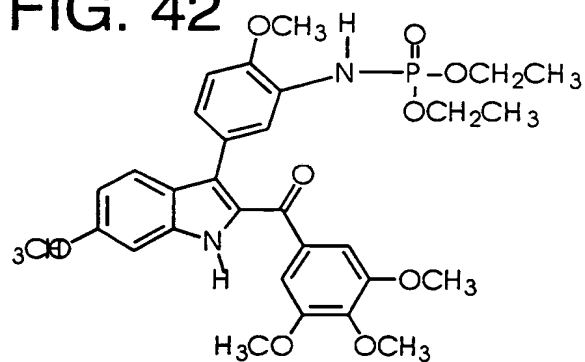


FIG. 43

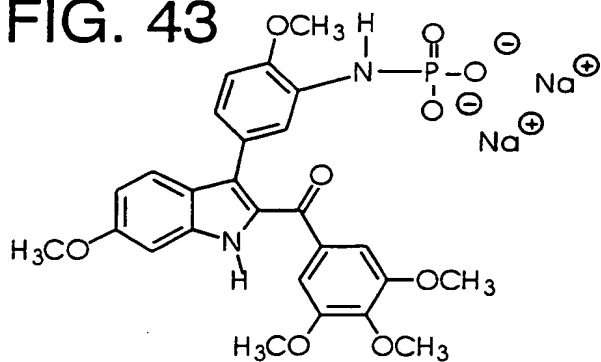


FIG. 44

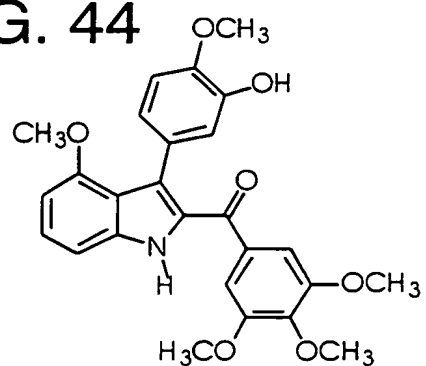


FIG. 45

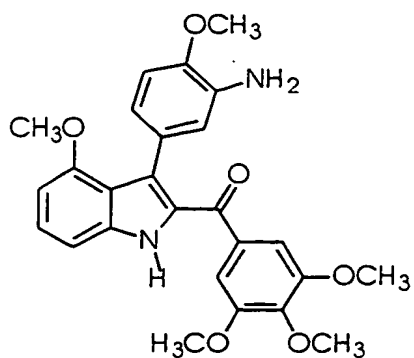


FIG. 46

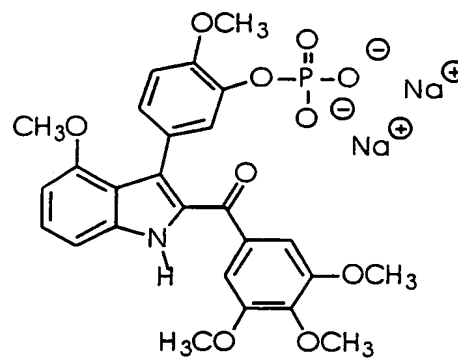


FIG. 47

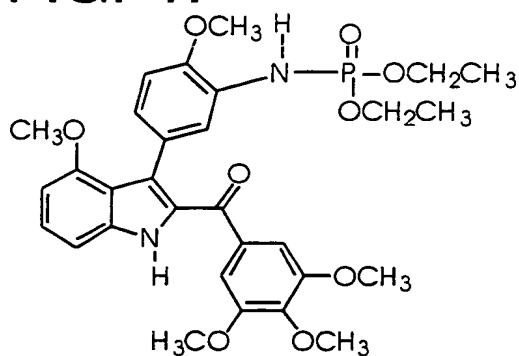


FIG. 48

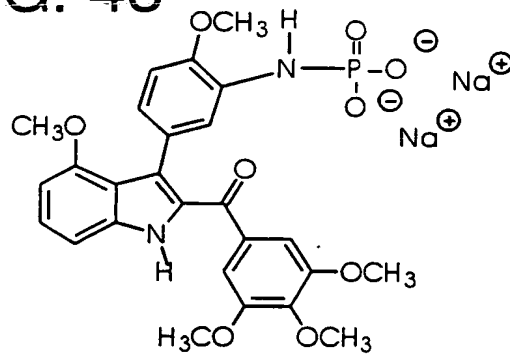


FIG. 49

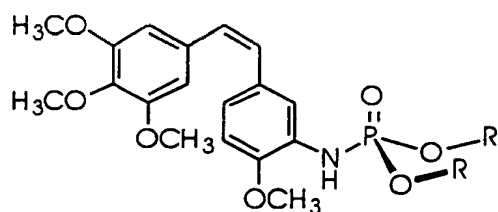


FIG. 50

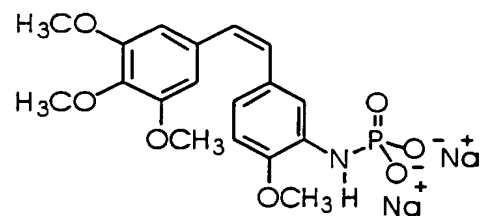
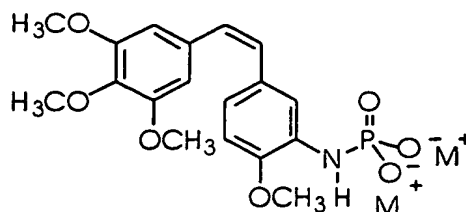


FIG. 51



(12) INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(19) World Intellectual Property Organization
International Bureau(43) International Publication Date
22 March 2001 (22.03.2001)

PCT

(10) International Publication Number
WO 01/19794 A3

- (51) International Patent Classification⁷: C07D 209/12, A61K 31/404, A61P 43/00, C07F 9/572, 9/24
- (74) Agent: HODGINS, Daniel, S.; Head, Johnson & Kachigian, 228 West 17th Place, Tulsa, OK 74119 (US).
- (21) International Application Number: PCT/US00/25408
- (81) Designated States (*national*): AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW.
- (22) International Filing Date:
15 September 2000 (15.09.2000)
- (25) Filing Language: English
- (26) Publication Language: English
- (84) Designated States (*regional*): ARIPO patent (GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).
- (30) Priority Data:
60/154,639 17 September 1999 (17.09.1999) US
- (71) Applicant (*for all designated States except US*): BAYLOR UNIVERSITY [US/US]; P.O. Box 97034, Waco, TX 76798-7034 (US).
- (72) Inventors; and
- (75) Inventors/Applicants (*for US only*): PINNEY, Kevin, G. [US/US]; 100 Russell Lane, Hewitt, TX 76643 (US). WANG, Feng [CN/CN]; 600 American Avenue, Apartment C304, King of Prussia, PA 19406 (US). DEL PILAR MEJIA, Maria [CO/US]; 9999 Linda Lane, Apartment GE, Des Plaines, IL 60016 (US).
- Published:
— with international search report
- (88) Date of publication of the international search report:
27 September 2001
- For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.*

(54) Title: INDOLE-CONTAINING AND COMBRETASTATIN-RELATED ANTI-MITOTIC AND ANTI-TUBULIN POLYMERIZATION AGENTS

(57) Abstract: Trimethoxyphenyl substituted indole ligands have been discovered which demonstrate impressive cytotoxicity as well as a remarkable ability to inhibit tubulin polymerization. Such compounds as well as related derivatives are excellent clinical candidates for the treatment of cancer in humans. In addition, certain of these ligands, as pro-drugs, may well prove to be tumor selective vascular targeting and destruction chemotherapeutic agents or to have anti-angiogenesis activity resulting in the selective prevention and/or destruction of tumor cell vasculature.

WO 01/19794 A3

INTERNATIONAL SEARCH REPORT

International Application No

PCT/US 00/25408

A. CLASSIFICATION OF SUBJECT MATTER

IPC 7 C07D209/12 31/404 A61P43/00 C07F9/24 C07F9/24

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 C07D C07F

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

CHEM ABS Data

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	WO 98 39323 A (BAYLOR UNIVERSITY) 11 September 1998 (1998-09-11) cited in the application claims ---	1-42
A	WO 99 34788 A (ARIZONA BOARD OF REGENTS) 15 July 1999 (1999-07-15) claims ---	37-42
P, X	WO 00 48606 A (OXIGENE, INC.) 24 August 2000 (2000-08-24) * table 2 and fig. 1B: compound X; fig. 1C: compound XVIII * -----	1-42



Further documents are listed in the continuation of box C.



Patent family members are listed in annex.

* Special categories of cited documents:

- *A* document defining the general state of the art which is not considered to be of particular relevance
- *E* earlier document but published on or after the international filing date
- *L* document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- *O* document referring to an oral disclosure, use, exhibition or other means
- *P* document published prior to the international filing date but later than the priority date claimed

T later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

X document of particular relevance: the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

Y document of particular relevance: the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

& document member of the same patent family

Date of the actual completion of the international search

24 April 2001

Date of mailing of the international search report

07/05/2001

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2
NL - 2280 HV Rijswijk
Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,
Fax: (+31-70) 340-3016

Authorized officer

Van Bijlen, H

INTERNATIONAL SEARCH REPORT

International Application No

PCT/US 00/25408

A. CLASSIFICATION OF SUBJECT MATTER

IPC 7 C07D209/12 A61P43/00 C07F9/572 C07F9/24

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IPC 7 C07D C07F

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Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

CHEM ABS Data

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A	WO 99 34788 A (ARIZONA BOARD OF REGENTS) 15 July 1999 (1999-07-15) claims	37-42
P, X	WO 00 48606 A (OXIGENE, INC.) 24 August 2000 (2000-08-24) * table 2 and fig. 1B: compound X; fig. 1C: compound XVIII *	1-42

☐

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- *O* document referring to an oral disclosure, use, exhibition or other means
- *P* document published prior to the international filing date but later than the priority date claimed

- *T* later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
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- *Y* document of particular relevance: the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.
- *Z* document member of the same patent family

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24 April 2001

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Name and mailing address of the ISA

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NL - 2280 HV Rijswijk
Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,
Fax: (+31-70) 340-3016

Authorized officer

Van Bijlen, H

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/US 00/25408

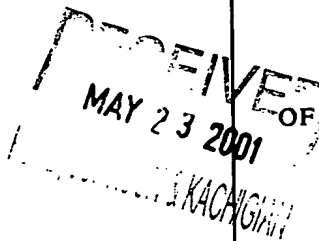
Patent document cited in search report		Publication date	Patent family member(s)		Publication date
WO 9839323	A	11-09-1998	US	5886025 A	23-03-1999
			AU	6688698 A	22-09-1998
			EP	0984954 A	15-03-2000
WO 9934788	A	15-07-1999	EP	1045689 A	25-10-2000
WO 0048606	A	24-08-2000	AU	3597300 A	04-09-2000

PATENT COOPERATION TREATY

From the
INTERNATIONAL PRELIMINARY EXAMINING AUTHORITY

To:

DANIEL S. HODGINS
HEAD, JOHNSON & KACHIGIAN
228 WEST 17TH PLACE
TULSA OK 74119



NOTIFICATION OF RECEIPT OF DEMAND BY COMPETENT INTERNATIONAL PRELIMINARY EXAMINING AUTHORITY

(PCT Rule 59.3(e) and 61.1(b), first sentence
and Administrative Instructions, Section 601(a))

Date of mailing
(day/month/year)

18 MAY 2001

Applicant's or agent's file reference
BAY515/00414

IMPORTANT NOTIFICATION

International application No.
PCT/US00/25408

International filing date (day/month/year)
15 SEP 00

Priority date (day/month/year)
17 SEP 99

Applicant

BAYLOR UNIVERSITY

1. The applicant is hereby **notified** that this International Preliminary Examining Authority considers the following date as the date of receipt of the demand for international preliminary examination of the international application:

16 April 2001 (16-04-01)

2. That date of receipt is:

- ☒ the actual date of receipt of the demand by this Authority (Rule 61.1(b)).
- ☐ the actual date of receipt of the demand on behalf of this Authority (Rule 59.3(e)).
- ☐ the date on which this Authority has, in response to the invitation to correct defects in the demand (Form PCT/IPEA/404), received the required corrections.

3. ☐ **ATTENTION:** That date of receipt is **AFTER** the expiration of 19 months from the priority date. Consequently, the election(s) made in the demand does (do) not have the effect of postponing the entry into the national phase until 30 months from the priority date (or later in some Offices) (Article 39(1)). Therefore, the acts for entry into the national phase must be performed within 20 months from the priority date (or later in some Offices) (Article 22). For details, see the *PCT Applicant's Guide*, Volume II.

- ☐ (If applicable) This notification confirms the information given by telephone, facsimile transmission or in person on:

4. Only where paragraph 3 applies, a copy of this notification has been sent to the International Bureau.

Name and mailing address of the IPEA/US
Assistant Commissioner for Patents
Box PCT
Washington, D.C. 20231
Facsimile No. Attn: IPEA/US

Authorized officer

Telephone No.

M. J. J.
703-308-6454

Form PCT/IPEA/402 (July 1998)

ATTORNEY

MAY 23 2001

ACTION DUE:
SETUP:

Noted

PATENT COOPERATION TREATY

PCT

INTERNATIONAL SEARCH REPORT

(PCT Article 18 and Rules 43 and 44)

Applicant's or agent's file reference BAY515/00414	FOR FURTHER ACTION see Notification of Transmittal of International Search Report (Form PCT/ISA/220) as well as, where applicable, item 5 below.	
International application No. PCT/US 00/ 25408	International filing date (day/month/year) 15/09/2000	(Earliest) Priority Date (day/month/year) 17/09/1999
Applicant BAYLOR UNIVERSITY		

This International Search Report has been prepared by this International Searching Authority and is transmitted to the applicant according to Article 18. A copy is being transmitted to the International Bureau.

This International Search Report consists of a total of 3 sheets.

☒ It is also accompanied by a copy of each prior art document cited in this report.

1. Basis of the report

- a. With regard to the **language**, the international search was carried out on the basis of the international application in the language in which it was filed, unless otherwise indicated under this item.

☐ the international search was carried out on the basis of a translation of the international application furnished to this Authority (Rule 23.1(b)).

- b. With regard to any **nucleotide and/or amino acid sequence** disclosed in the international application, the international search was carried out on the basis of the sequence listing :

☐ contained in the international application in written form.

☐ filed together with the international application in computer readable form.

☐ furnished subsequently to this Authority in written form.

☐ furnished subsequently to this Authority in computer readable form.

☐ the statement that the subsequently furnished written sequence listing does not go beyond the disclosure in the international application as filed has been furnished.

☐ the statement that the information recorded in computer readable form is identical to the written sequence listing has been furnished

2. ☒ **Certain claims were found unsearchable** (See Box I).

3. ☐ **Unity of invention is lacking** (see Box II).

4. With regard to the title,

☒ the text is approved as submitted by the applicant.

☐ the text has been established by this Authority to read as follows:

5. With regard to the abstract,

☒ the text is approved as submitted by the applicant.

☐ the text has been established, according to Rule 38.2(b), by this Authority as it appears in Box III. The applicant may, within one month from the date of mailing of this international search report, submit comments to this Authority.

6. The figure of the drawings to be published with the abstract is Figure No.

☐ as suggested by the applicant.

☐ because the applicant failed to suggest a figure.

☐ because this figure better characterizes the invention.

☐ None of the figures.

INTERNATIONAL SEARCH REPORT

International Application No

PCT/US 00/25408

A. CLASSIFICATION OF SUBJECT MATTER

IPC 7 C07D209/12 A61K31/404 A61P43/00 C07F9/572 C07F9/24

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 C07D C07F

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

CHEM ABS Data

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
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P, X	WO 00 48606 A (OXIGENE, INC.) 24 August 2000 (2000-08-24) * table 2 and fig. 1B: compound X; fig. 1C: compound XVIII *	1-42

☐ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

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- *E* earlier document but published on or after the international filing date
- *L* document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
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- *X* document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
- *Y* document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.
- *&* document member of the same patent family

Date of the actual completion of the international search

24 April 2001

Date of mailing of the international search report

07/05/2001

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2
NL - 2280 HV Rijswijk
Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,
Fax: (+31-70) 340-3016

Authorized officer

Van Bijlen, H

INTERNATIONAL SEARCH REPORT

International application No.
PCT/US 00/25408

Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☒ Claims Nos.:
because they relate to subject matter not required to be searched by this Authority, namely:

Although claims 43 and 44 are directed to a method of treatment of the human/animal body, the search has been carried out and based on the alleged effects of the compound/composition.
2. ☐ Claims Nos.:
because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:
3. ☐ Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. ☐ As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest.
- ☐ No protest accompanied the payment of additional search fees.

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/US 00/25408

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
WO 9839323 A	11-09-1998	US 5886025 A	23-03-1999
		AU 6688698 A	22-09-1998
		EP 0984954 A	15-03-2000
WO 9934788 A	15-07-1999	EP 1045689 A	25-10-2000
WO 0048606 A	24-08-2000	AU 3597300 A	04-09-2000

(12) INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(19) World Intellectual Property Organization
International Bureau(43) International Publication Date
22 March 2001 (22.03.2001)

PCT

(10) International Publication Number
WO 01/19794 A3

- (51) International Patent Classification⁷: C07D 209/12, A61K 31/404, A61P 43/00, C07F 9/572, 9/24
- (74) Agent: HODGINS, Daniel, S.; Head, Johnson & Kachigian, 228 West 17th Place, Tulsa, OK 74119 (US).
- (21) International Application Number: PCT/US00/25408
- (81) Designated States (*national*): AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW.
- (22) International Filing Date:
15 September 2000 (15.09.2000)
- (25) Filing Language: English
- (26) Publication Language: English
- (30) Priority Data:
60/154,639 17 September 1999 (17.09.1999) US
- (84) Designated States (*regional*): ARIPO patent (GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).
- (71) Applicant (*for all designated States except US*): BAYLOR UNIVERSITY [US/US]; P.O. Box 97034, Waco, TX 76798-7034 (US).
- Published:
— with international search report
- (72) Inventors; and
- (75) Inventors/Applicants (*for US only*): PINNEY, Kevin, G. [US/US]; 100 Russell Lane, Hewitt, TX 76643 (US). WANG, Feng [CN/CN]; 600 American Avenue, Apartment C304, King of Prussia, PA 19406 (US). DEL PILAR MEJIA, Maria [CO/US]; 9999 Linda Lane, Apartment GE, Des Plaines, IL 60016 (US).
- (88) Date of publication of the international search report:
27 September 2001
- For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.*

(54) Title: INDOLE-CONTAINING AND COMBRETASTATIN-RELATED ANTI-MITOTIC AND ANTI-TUBULIN POLYMERIZATION AGENTS

(57) Abstract: Trimethoxyphenyl substituted indole ligands have been discovered which demonstrate impressive cytotoxicity as well as a remarkable ability to inhibit tubulin polymerization. Such compounds as well as related derivatives are excellent clinical candidates for the treatment of cancer in humans. In addition, certain of these ligands, as pro-drugs, may well prove to be tumor selective vascular targeting and destruction chemotherapeutic agents or to have anti-angiogenesis activity resulting in the selective prevention and/or destruction of tumor cell vasculature.

WO 01/19794 A3

PATENT COOPERATION TREATY

PCT

NOTIFICATION OF THE RECORDING
OF A CHANGE(PCT Rule 92bis.1 and
Administrative Instructions, Section 422)

From the INTERNATIONAL BUREAU

To:

HODGINS, Daniel, S.
Head, Johnson & Kachigian
228 West 17th Place
Tulsa, OK 74119
ETATS-UNIS D'AMERIQUE

Date of mailing (day/month/year) 23 January 2001 (23.01.01)	IMPORTANT NOTIFICATION International filing date (day/month/year) 15 September 2000 (15.09.00)
Applicant's or agent's file reference BAY515/00414	
International application No. PCT/US00/25408	

1. The following indications appeared on record concerning: <input checked="" type="checkbox"/> the applicant <input checked="" type="checkbox"/> the inventor <input type="checkbox"/> the agent <input type="checkbox"/> the common representative		
Name and Address DEL PILAR MEJIA, Maria 1239 Emerson Street Apartment 2 Evanston, IL 60201 United States of America	State of Nationality CO	State of Residence US
	Telephone No.	
	Facsimile No.	
	Teleprinter No.	
2. The International Bureau hereby notifies the applicant that the following change has been recorded concerning: <input type="checkbox"/> the person <input type="checkbox"/> the name <input checked="" type="checkbox"/> the address <input type="checkbox"/> the nationality <input type="checkbox"/> the residence		
Name and Address DEL PILAR MEJIA, Maria 9999 Linda Lane Apartment GE Des Plaines, IL 60016 United States of America	State of Nationality	State of Residence
	Telephone No.	
	Facsimile No.	
	Teleprinter No.	
3. Further observations, if necessary:		
4. A copy of this notification has been sent to: <input checked="" type="checkbox"/> the receiving Office <input checked="" type="checkbox"/> the designated Offices concerned <input checked="" type="checkbox"/> the International Searching Authority <input type="checkbox"/> the elected Offices concerned <input type="checkbox"/> the International Preliminary Examining Authority <input type="checkbox"/> other:		

The International Bureau of WIPO 34, chemin des Colombettes 1211 Geneva 20, Switzerland Facsimile No.: (41-22) 740.14.35	Authorized officer J. Leitao Telephone No.: (41-22) 338.83.38
---	---

PATENT COOPERATION TREATY

PCT

NOTIFICATION OF ELECTION

(PCT Rule 61.2)

From the INTERNATIONAL BUREAU

To:

Commissioner
 US Department of Commerce
 United States Patent and Trademark
 Office, PCT
 2011 South Clark Place Room
 CP2/5C24
 Arlington, VA 22202
 ETATS-UNIS D'AMERIQUE

in its capacity as elected Office

Date of mailing (day/month/year) 13 July 2001 (13.07.01)	
International application No. PCT/US00/25408	Applicant's or agent's file reference BAY515/00414
International filing date (day/month/year) 15 September 2000 (15.09.00)	Priority date (day/month/year) 17 September 1999 (17.09.99)
Applicant PINNEY, Kevin, G. et al	

1. The designated Office is hereby notified of its election made:

☒ in the demand filed with the International Preliminary Examining Authority on:

16 April 2001 (16.04.01)

☐ in a notice effecting later election filed with the International Bureau on:

2. The election
- ☒
- was

☐ was not

made before the expiration of 19 months from the priority date or, where Rule 32 applies, within the time limit under Rule 32.2(b).

The International Bureau of WIPO 34, chemin des Colombettes 1211 Geneva 20, Switzerland Facsimile No.: (41-22) 740.14.35	Authorized officer Claudio Borton Telephone No.: (41-22) 338.83.38
---	--

PATENT COOPERATION TREATY

PCT

NOTIFICATION OF THE RECORDING
OF A CHANGE(PCT Rule 92bis.1 and
Administrative Instructions, Section 422)

From the INTERNATIONAL BUREAU

To:

HODGINS, Daniel, S.
Jackson Walker L.L.P.
Suite 2100
112 E. Pecan
San Antonio, TX 78205
ETATS-UNIS D'AMERIQUEDate of mailing (day/month/year)
18 février 2002 (18.02.02)Applicant's or agent's file reference
P 208614.9PCT

IMPORTANT NOTIFICATION

International application No.
PCT/US00/25408International filing date (day/month/year)
15 septembre 2000 (15.09.00)

1. The following indications appeared on record concerning:

☐ the applicant ☐ the inventor ☒ the agent ☐ the common representative

Name and Address

HODGINS, Daniel, S.
Head, Johnson & Kachigian
228 West 17th Place
Tulsa, OK 74119
United States of America

State of Nationality

State of Residence

Telephone No.

1 918 587 2000

Facsimile No.

1 918 587 5603

Teleprinter No.

2. The International Bureau hereby notifies the applicant that the following change has been recorded concerning:

☐ the person ☐ the name ☒ the address ☐ the nationality ☐ the residence

Name and Address

HODGINS, Daniel, S.
Jackson Walker L.L.P.
Suite 2100
112 E. Pecan
San Antonio, TX 78205
United States of America

State of Nationality

State of Residence

Telephone No.

(210) 978-7700

Facsimile No.

(210) 978-7790

Teleprinter No.

3. Further observations, if necessary:

The agent's file reference has been changed accordingly.

4. A copy of this notification has been sent to:

☒ the receiving Office ☐ the designated Offices concerned
☐ the International Searching Authority ☒ the elected Offices concerned
☒ the International Preliminary Examining Authority ☐ other:The International Bureau of WIPO
34, chemin des Colombettes
1211 Geneva 20, Switzerland

Authorized officer

François BAECHLER

Facsimile No.: (41-22) 740.14.35

Telephone No.: (41-22) 338.83.38

PATENT COOPERATION TREATY

PCT

INTERNATIONAL SEARCH REPORT

(PCT Article 18 and Rules 43 and 44)

Applicant's or agent's file reference BAY515/00414	FOR FURTHER ACTION see Notification of Transmittal of International Search Report (Form PCT/ISA/220) as well as, where applicable, item 5 below.	
International application No. PCT/US 00/ 25408	International filing date (day/month/year) 15/09/2000	(Earliest) Priority Date (day/month/year) 17/09/1999
Applicant BAYLOR UNIVERSITY		

This International Search Report has been prepared by this International Searching Authority and is transmitted to the applicant according to Article 18. A copy is being transmitted to the International Bureau.

This International Search Report consists of a total of 3 sheets.

☒ it is also accompanied by a copy of each prior art document cited in this report.

1. Basis of the report

- a. With regard to the **language**, the international search was carried out on the basis of the international application in the language in which it was filed, unless otherwise indicated under this item.

☐ the international search was carried out on the basis of a translation of the international application furnished to this Authority (Rule 23.1(b)).

- b. With regard to any **nucleotide and/or amino acid sequence** disclosed in the international application, the international search was carried out on the basis of the sequence listing :

☐ contained in the international application in written form.

☐ filed together with the international application in computer readable form.

☐ furnished subsequently to this Authority in written form.

☐ furnished subsequently to this Authority in computer readable form.

☐ the statement that the subsequently furnished written sequence listing does not go beyond the disclosure in the international application as filed has been furnished.

☐ the statement that the information recorded in computer readable form is identical to the written sequence listing has been furnished

2. ☒ **Certain claims were found unsearchable** (See Box I).

3. ☐ **Unity of invention is lacking** (see Box II).

4. With regard to the **title**,

☒ the text is approved as submitted by the applicant.

☐ the text has been established by this Authority to read as follows:

5. With regard to the **abstract**,

☒ the text is approved as submitted by the applicant.

☐ the text has been established, according to Rule 38.2(b), by this Authority as it appears in Box III. The applicant may, within one month from the date of mailing of this international search report, submit comments to this Authority.

6. The figure of the **drawings** to be published with the abstract is Figure No. _____

☐ as suggested by the applicant.

☐ because the applicant failed to suggest a figure.

☐ because this figure better characterizes the invention.

☐ None of the figures.

INTERNATIONAL SEARCH REPORT

International Application No

PC 00/25408

A. CLASSIFICATION OF SUBJECT MATTER

IPC 7 C07D209/12 A61K31/404 A61P43/00 C07F9/572 C07F9/24

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 C07D C07F

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

CHEM ABS Data

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	WO 98 39323 A (BAYLOR UNIVERSITY) 11 September 1998 (1998-09-11) cited in the application claims ---	1-42
A	WO 99 34788 A (ARIZONA BOARD OF REGENTS) 15 July 1999 (1999-07-15) claims ---	37-42
P, X	WO 00 48606 A (OXIGENE, INC.) 24 August 2000 (2000-08-24) * table 2 and fig. 1B: compound X; fig. 1C: compound XVIII * -----	1-42

☐

Further documents are listed in the continuation of box C.

☒

Patent family members are listed in annex.

* Special categories of cited documents:

- *A* document defining the general state of the art which is not considered to be of particular relevance
- *E* earlier document but published on or after the international filing date
- *L* document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- *O* document referring to an oral disclosure, use, exhibition or other means
- *P* document published prior to the international filing date but later than the priority date claimed

- *T* later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
- *X* document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
- *Y* document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.
- * & * document member of the same patent family

Date of the actual completion of the international search

24 April 2001

Date of mailing of the international search report

07/05/2001

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2
NL - 2280 HV Rijswijk
Tel (+31-70) 340-2040, Tx. 31 651 epo nl,
Fax: (+31-70) 340-3016

Authorized officer

Van Bijlen, H

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/US 00/25408

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
WO 9839323 A	11-09-1998	US 5886025 A	23-03-1999
		AU 6688698 A	22-09-1998
		EP 0984954 A	15-03-2000
WO 9934788 A	15-07-1999	EP 1045689 A	25-10-2000
WO 0048606 A	24-08-2000	AU 3597300 A	04-09-2000